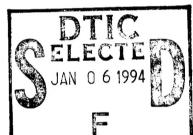


FINAL REPORT FOR SOURCE TEST MEASUREMENT OF NITROGEN OXIDES, SULFUR DIOXIDE, CARBON MONOXIDE, VOC, AND PM10 EMISSIONS ON GAS TURBINE #2 AT ONIZUKA AIR FORCE BASE SUNNYVALE, CALIFORNIA

S. HUGH BROWN

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FEBRUARY 1994

Final Contractor Report for Period December 1993 - January 1994

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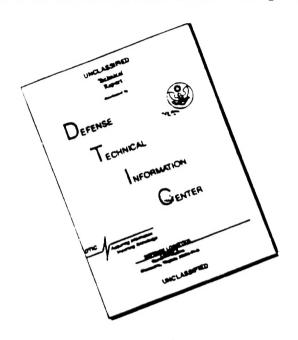
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FOREWORD

This report was prepared by Pacific Environmental Services, Inc. (PES) using due and ordinary care and in accordance with the scope of work identified in the Professional Services Agreement, Purchase Order, or other written/verbal request.

Any measured contaminants and their concentrations detected or identified in this study are for the sample(s) obtained or for the observations made at the time(s) and location(s) referenced in the report and may not represent other times, locations, or contaminants and their concentrations.

Except for the tests and observations conducted by PES, no attempt was made to check for compliance of present or past owners or operators of the equipment, plant, or site with federal, state, or local laws and regulations.

The information provided in this report, including any drawings and specifications, was prepared solely for the use of the identified client and any use by any other party shall be at their own risk.

The project work was conducted by Steven M. Hernandez, Robert T. Nguyen, and S. Hugh Brown under the direction of S. Hugh Brown.

Approved:

S. Hugh Brown, Director Air Quality Testing

5 High Brown

A-1

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INTRODUCTION

Onizuka Air Force Base currently has twelve gas turbine generators which supply electric power for the entire facility. Heat recovery boilers are connected to each generator. Current plans are for the power plant to curtail its operation. Under current Bay Area Air Quality Management District (BAAQMD) rules and regulations, emissions credits can be banked and sold. Source testing is required to quantify the baseline emissions for each constituent potentially reduced.

Based on information obtained from the BAAQMD, source testing was conducted for nitrogen oxides (NO_x), sulfur dioxide (SO₂), carbon monoxide (CO), volatile organic compounds (VOC), and particulate matter less than 10 microns (PM10).

Pacific Environmental Services (PES), a participant in CARB's Independent Contractor Program, was hired by the Air Force to perform the required source testing and data reduction. PES qualifies as an independent testing laboratory (no conflict of interest). The source testing was conducted by S. Hugh Brown, Steven Hernandez, and Robert Nguyen of PES on December 3, 1993 and January 12, 1994.

EQUIPMENT AND PROCESS DESCRIPTION

All of the gas turbine generators were identical 750 kVA units that were manufactured by Solar and each consumed a maximum of 130 therms/hour of natural gas. Gas turbine #2 was equipped with sampling ports and a work platform and was used as a demonstration unit for all twelve generators. The turbine exhaust gases were routed under the turbine platform and over to the heat recovery boiler adjacent to the turbine on the same platform. The gases entered the boiler through a diverter/bypass damper at the base of the exhaust stack. When the damper was in the bypass position, the gases were routed straight up the stack to atmosphere. When the damper was in the closed position, the gases were routed horizontally through the boiler exiting the top and rejoining the stack just above the bypass damper. The location of the turbines on the base is shown in Figure 2.1. The gas turbine and heat recovery boiler are depicted in Figure 2.2. A copy of the Permit to Operate issued by the BAAQMD is located in Appendix A.

The source testing was conducted on gas turbine #2 at a constant load of about 570 KW with the heat recovery boiler damper set at 75% closed (25% bypass).

Figure 2.1
Plant Layout And Equipment Location

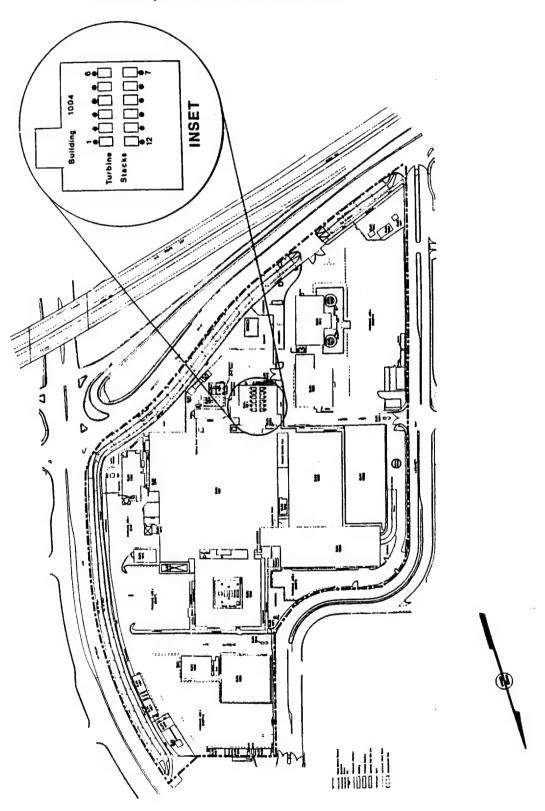
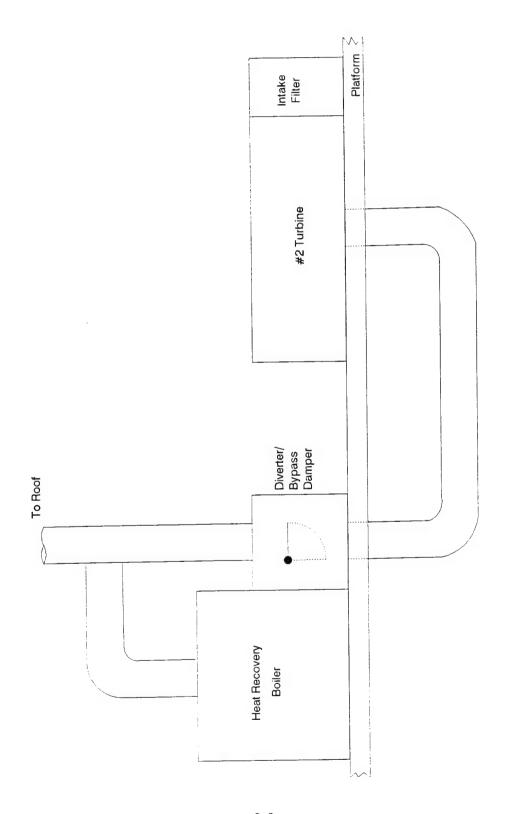


Figure 2.2 Schematic - Turbine/Heat Recovery Boiler Number 2



TESTING METHODOLOGY

The approximate sampling port locations are shown on Figure 3.1. The number of traverse points required (4 on each of two diameters, 90 degrees apart) and their locations were derived from CARB Method 1. The sampling was conducted on the exhaust of turbine #2 at roof level during block load conditions. Triplicate tests were conducted for each constituent determined.

PM10

The PM10 emissions of the exhaust were determined by EPA Method 201A. The sampling train is shown in Figure 3.2 and consisted of a stainless steel nozzle, a 10-micron cut-off cyclone and an in-stack stainless steel filter holder manufactured by Andersen, a 48-inch glass probe, a 10-foot Teflon hose from the probe to the first impinger, two Greenburg-Smith impingers each charged with 100 mls of distilled water, an empty impinger, an impinger filled with silica gel, a 30-foot umbilical line, a vacuum pump, a dry gas meter and a calibrated orifice connected to an oil inclined manometer. Glass fiber filters were used in the filter holders. Each sample was collected isokinetically at a fixed sampling rate and the number of minutes sampled at each traverse point was determined by the ratio of the point velocity to the average velocity. The velocity and temperature were measured at each traverse point as it was sampled. Field data and calculation sheets are included in Appendix B.

The volume of the impinger solution and the weight of the silica gel were recorded before and after the tests in order to obtain the moisture content of the stack gas stream. All sample volumes and weights were recorded immediately on sample recovery sheets (Appendix B) during charging and sample recovery. Leak checks were performed before and after each test. The post-test leak check was performed after removing the cut-off cyclone so as not to disturb the particle catch.

After the test, the contents of the nozzle and cut-off cyclone (PM10+) were recovered by rinsing three times with acetone. The washings were placed in a 125-milliliter polyethylene container. The filter was placed in a plastic cassette. The contents of the cyclone exit and filter holder front (PM10) were also recovered by rinsing three times with acetone. The washings were placed in a separate 125-milliliter polyethylene container. The contents of each impinger set was placed in a 1000-milliliter polyethylene container. The sampling train was then rinsed from the 3rd impinger to the nozzle with the distilled water and the rinse was added to the sample. The impinger solution was saved but not analyzed. Disposable vinyl gloves were worn during sample retrieval to help prevent contamination.

Figure 3.1 Sampling Location - Turbine Number 2

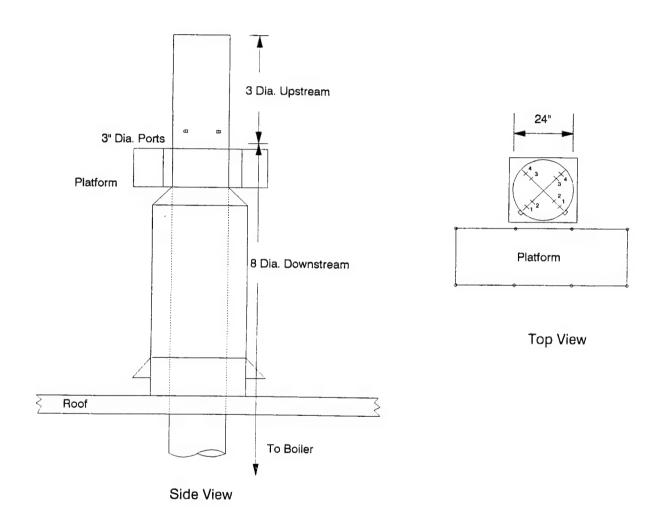
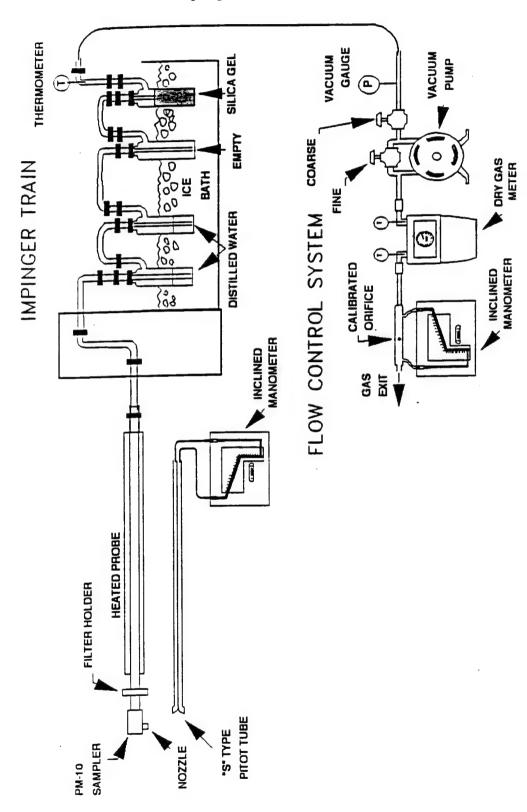


Figure 3.2
PM10 Sampling Train - EPA Method 201A



The particulate fractions were evaporated to dryness at 100 degrees Centigrade and desiccated to a constant weight along with the sample filter.

CONTINUOUS MONITORING

The turbine/boiler exhaust was monitored for VOC, CO, CO₂, O₂ and NO_x by CARB Method 100. Each test run was 60 minutes long. Rosemount Analytical Model 880 Infrared Analyzers were used to determine the CO and CO₂ concentrations. A Rosemount Analytical Model 755R Paramagnetic Analyzer was used to determine the oxygen concentration, and a Thermo Electron Model 10 Chemilumine-scent Analyzer was used to determine the NO_x concentration. The output of the analyzers was linearized by the manufacturers. Table 3.1 lists the instrument specifications. Table 3.2 lists the calibration gases that were used specific to this job.

The continuous monitoring train for the above gases is shown in Figure 3.3 and consisted of a 3/8-inch stainless steel sampling probe, a 3/8-inch heated Teflon sampling line, a sample refrigeration/pump system, a glass fiber filter in a 47-millimeter stainless steel holder, and a sample distribution manifold. The distribution manifold was equipped with a series of 3-way valves with flow meters (rotometer style). One flow meter acted as a bypass, and the others were connected to the individual analyzers. The output of the analyzers was logged by a Yokogawa Model HR2400 multi-channel recorder and a Rustrak Ranger II data logger.

The NO_x analyzer was operated on a range of 0-50 ppmv with span gases at 10.4, 23.5, and 44.2 ppmv. The CO analyzer was operated on a range of 0-100 ppmv with span gases at 50 and 75 ppmv. The O_2 analyzer was operated on a range of 0-25 per cent with span gases at 11.0 and 19.0 per cent. The CO_2 analyzer was operated on a range of 0-5 per cent with span gases at about 2.0 and 4.0 per cent. Prior to the source tests, the suction side of the monitoring system was leak-checked at a full vacuum (greater than 20 inches mercury).

A bias check was made on each analyzer by comparing the response between the span gas introduced at the sample line tip and the span gas introduced directly to the analyzer to ensure a differential of less than 5 per cent. The analyzers were spanned before and after each test run with NIST traceable calibration gases from Scott Specialty, and with zero grade nitrogen.

The total hydrocarbons were determined on January 12, 1994 by using a JUM Model VE-7 total hydrocarbon analyzer that utilized a heated Teflon sample line and a flame ionization detector mounted in a heated oven. The sample line was maintained at 250 degrees Fahrenheit, and the oven was maintained at 180 degrees Centigrade. Since the gas stream was expected to have a low concentration of

Table 3.1 Continuous Monitoring Specifications

Table 3.1 - Continued

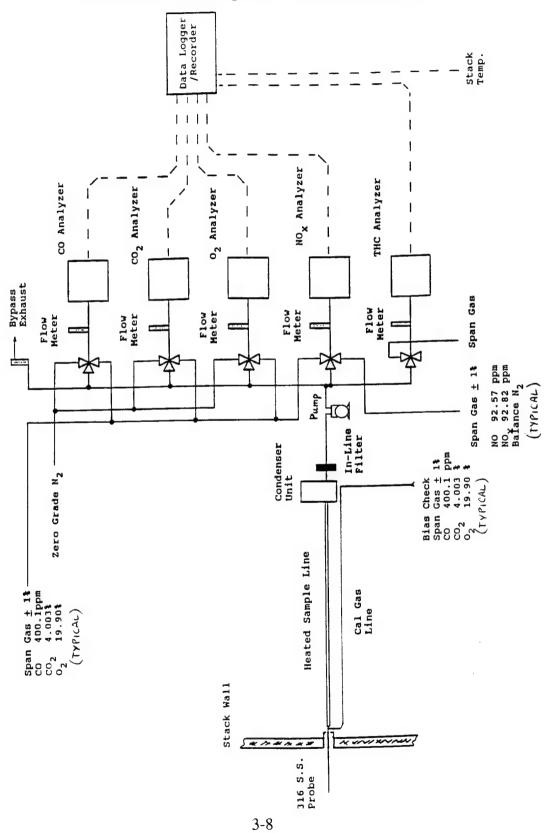
VOC Heated Hydrocarbon Analyzer Analyzer	- J.U.M. Engineeri	ng Model VE	-7 FID		
Response Time	0-95% in less than	1.2 Seconds			
Zero Drift	± 1% of Full Sca	± 1% of Full Scale in 24 Hrs			
Span Drift	+ 1% of Full scal	± 1% of Full scale in 24 Hrs			
Linearity	± 1% of Full Scale				
Accuracy	Derived from the Calibration Gas + 1% gas was used				
Sensitivity	1 ppb				
Range Change Consistency	Less than 1% Full Scale				
Oxygen Synergism	Less than 2%				
Output	0-10.0 Vdc Scaled	l :			
	$egin{array}{c} R_1 \ R_2 \ R_3 \ R_4 \ R_5 \end{array}$	0-10 0-100 0-1,000 0-10,000 0-100,000	ppm as C ₃ ppm as C ₃ ppm as C ₃ ppm as C ₃ ppm as C ₃		
Sample Flow Rate	3 Liters/Minute				

Table 3.2 Calibration Gases

Gas Composition	Use	Cylinder Ser. / No.	Certified Accuracy	Analysis Date
Nitrogen	Zero Gas	AAL / 4542	Zero Grade	N/A
11.0% O ₂ 50.0 ppmv CO 11.0% CO ₂ Bal N ₂	Span Gas	1L / 2572	± 1% ± 1% ± 1%	04/08/93
19.0% O ₂ 75.0 ppm CO 18% CO ₂ Bal N ₂	Span Gas	ALMO / 04292	<u>+</u> 1%	04/29/93
10.36 ppm NO 10.39 ppm NO _x Bal N ₂	Span Gas	ALM / 010841	<u>+</u> 1%*	12/02/93 exp 12/95
22.90 ppm NO 23.46 ppm NO_X Bal N_2	Span Gas	ALM / 034155	<u>+</u> 1%*	10/06/93 exp 10/95
43.83 ppm NO 44.17 ppm NO _x Bal N ₂	Span Gas	ALM / 027046	<u>+</u> 1%*	12/23/93 exp 12/95
2.0% CO	Span Gas	ALM / 008830	<u>+</u> 1%*	09/17/93
4.0 CO	Span Gas	ALM / 033923	<u>+</u> 1%*	09/16/93

^{*} EPA Protocol 1 gas.

Figure 3.3 Continuous Monitoring Train - CARB Method 100



hydrocarbons, the instrument was used on a 0-50 ppmv range and standardized with 20 and 40 ppmv propane calibration gases. Each test run was about 50 minutes long.

In order to determine the methane concentration of the stack gas for correcting the total hydrocarbon monitoring data and as a back-up total hydrocarbon analysis, an integrated sample of the flue gas was collected for about 40 minutes concurrently with each monitoring run by using EPA Method 25 modified to eliminate the condensate trap. The sampling train is shown in Figure 3.4 and consisted of a stainless steel sampling probe connected through a flow control device (micro orifice disc meter) to an evacuated 12-liter stainless steel cylinder. The orifice meter reading and the cylinder gage vacuum were recorded at 10-minute intervals during the sampling period on field data sheets (see Appendix B).

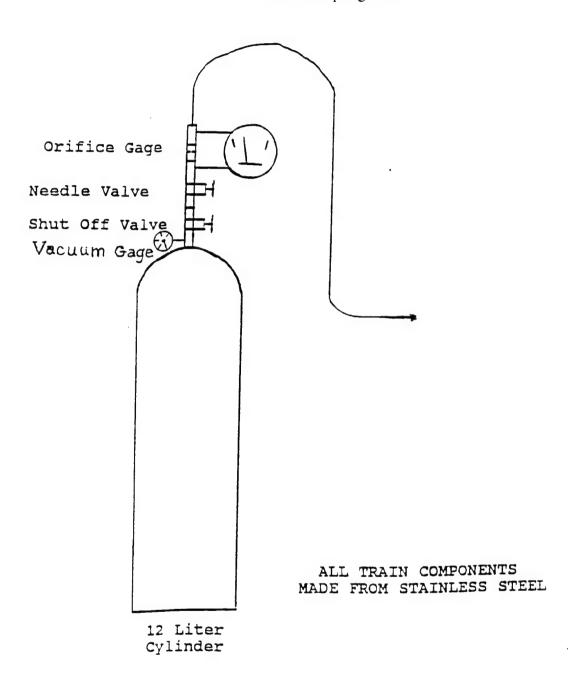
The integrated samples were analyzed for carbon monoxide (CO), methane (CH_4) , carbon dioxide (CO_2) , oxygen (O_2) , and nonmethane hydrocarbons (as C_1) by Truesdail Laboratories, in Tustin, CA according to EPA Method 25. A sample submittal/chain of custody sheet was completed and is included in Appendix B along with the field data and calculation sheets. The analytical procedures performed at Truesdail are detailed in Appendix C along with the laboratory report.

Emission factors for the gases were calculated based on the natural gas usage. A mass balance calculation was performed on the entire power plant. The natural gas usage of all turbines is routed through a gas meter. The gas usage was monitored for first test day and related to the number of turbines operating and the kilowatt load for each.

Sulfur Dioxide

Sulfur dioxide emissions were expected to be very low. Since the source of sulfur is the odorant in the natural gas, the sulfur dioxide emissions were reported in pounds per therm (100 CF of natural gas) by using the sulfur content of the fuel (from the gas supplier).

Figure 3.4 VOC Sampling Train



NOTE: Samples usually collected in duplicate or triplicate concurrently

RESULTS

Calculations were made from the field data sheets and lab analyses to determine sample volume, molecular weight, velocities, flow rates, component concentrations, and micron cut-off size for the tests.

PM10

Analyses of PM10 emissions for each test run on the turbine included determinations for total solid particulate matter (PM). Condensables were not included in the PM10 results. Although some weight was recorded for each PM10 fraction, 1 milligram was chosen as a level of significance in order to account for fluctuations in the balance weights and for minute amounts of grease contamination from the threads of the PM10 sampling head. Results of the tests are summarized in Table 4.1. No significant PM10 or particulate matter was detected.

Gases

The nitrogen oxide and carbon monoxide emissions were 3.1 and 3.5 pounds per hour, respectively. The emission factors were reported as pounds per therm (100 CF) as indicated on the natural gas meter supplying the power plant. The gas meter had two indicators: one mechanical and one digital. The larger volume reading (digital) was assumed to be the corrected reading and this value was used for all of the emission factors calculations. However, this reading appears to have some unknown multiplier because the therm usage is too low for the known consumption of the gas turbines. For the purposes of calculating any periodic emissions, the emission factors should be used against the digital reading as indicated at the gas meter. Results are reported in Table 4.1.

The continuous monitoring for total hydrocarbons was unexpectedly variable and high (10 to 35 ppmv as C₃) and indicated that the sample line, although heated, was contaminated from previous testing which involved very heavy organics. As a back-up, the integrated sample cylinders that were sampled concurrently and analyzed for methane were also analyzed for total nonmethane hydrocarbons and the resulting average value (5 ppmv) was used to calculate the VOC emission factor. Results are reported in Table 4.2.

The sulfur dioxide emission factor was calculated from information supplied by PG&E in San Francisco, CA which supplies the natural gas to the base. The sulfur dioxide emission factor used by PG&E is 0.001 pounds per million BTU which

Table 4.1

Turbine Number 2 Exhaust - PM10 Continuous Emission Monitoring

Description	Run #1	Run #2	Run #3	Average
Sampling Date Sample Number Turbine Load, KW Waste Heat Damper, % Gas Usage, therms/hr	12/03/93 OAFB-1 570 75 12.4*	12/03/93 OAFB-2 570 75 12.4*	12/03/93 OAFB-3 558 75 12.2*	
Flue Gas Temperature, °F Velocity, ft/sec Flow Rate, ACFM Flow Rate, DSCFM Moisture, % v/v	450 117.4 22,120 12,300 4.7	457 117.1 22,060 12,200 4.5	460 115.8 21,820 12,030 4.5	
PM10 Sample Start Sample Stop Sampling Time, min Sample volume, DSCF Cutoff, microns Collection, g Concentration, g/DSCF Emissions, lbs/hr Factor, lbs/therm	07:19 09:22 116.1 90.1 10.0 <0.001 <0.0002 <0.002 <0.002*	10:30 12:29 115.2 90.3 10.0 <0.001 <0.0002 <0.02 <0.002*	13:54 15:51 113.7 89.1 10.0 <0.001 <0.0002 <0.002 <0.002*	<0.0002 <0.02 <0.002
Total PM Collection, g Concentration, g/DSCF Emissions, lbs/hr	<0.001 <0.0004 <0.04	<0.001 <0.0004 <0.04	<0.001 <0.0004 <0.04	<0.0004 <0.04

Table 4.1 - Continued

Description	Run #1	Run #2	Run #3	Average
Continuous Monitoring - Gases Sample Start Sample Stop Concentration, % v/v Carbon Dioxide Oxygen Concentration, ppmv Nitrogen Oxides Carbon Monoxide Emissions, lbs/hr	09:56 10:56 2.1 17.6 34 66	11:51 12:51 2.2 17.5 36 66	Run #3 13:46 14:46 2.2 17.5 37 66	2.2 17.5 36 66
Nitrogen Oxides	3.0	3.1	3.1	3.1
Carbon Monoxide	3.5	3.5	3.5	3.5
Factor, lbs/therm Nitrogen Oxides Carbon Monoxide	0.24*	0.25*	0.25*	0.25*
	0.28*	0.28*	0.29*	0.28*

^{*} Therms as indicated on digital readout of gas meter - appears to have some unknown multiplier.

Table 4.2 Turbine Number 2 Exhaust - Hydrocarbon Emissions

Description	Run #1	Run #2	Run #3	Average
Sampling Date Turbine Load, KW Waste Heat Damper, % Gas Usage, therms/hr	01/12/94 550 75 12.0*	01/12/94 550 75 12.0*	01/12/94 550 75 12.0*	
Flue Gas Temperature, °F Velocity, ft/sec Flow Rate, ACFM Flow Rate, DSCFM Moisture, % v/v	441 103.2 19,450 10,930 4.6**	444 103.2 19,450 10,930 4.6**	445 103.2 19,450 10,930 4.6**	
Hydrocarbon Samples Sample Start Sample Stop Sampling Time, min Concentration, % v/v Oxygen (O ₂) Concentration, ppmv C ₁ Carbon Dioxide (CO ₂) Carbon Monoxide (CO) Methane (CH ₄) Total HC (w/o CH ₄) Emission Rate, lbs/hr C ₁ Total HC (w/o CH ₄) Emissions Factor, lbs/therm Total HC (w/o CH ₄)	10:56 11:36 40 17.9 18,610 60 7 <4 <0.04 <0.003*	12:18 12:58 40 18.2 18,590 58 6 11 0.22 0.018*	13:38 14:18 40 18.1 18,700 57 6 <4 <0.04 <0.003*	18.1 18,630 58 6 5

^{*} Therms as indicated on digital readout of gas meter - appears to have some unknown multiplier.
* From PM10 testing - 12/3/93.

is equivalent to a concentration of 6 ppmv in the natural gas as hydrogen sulfide (H_2S) . Results are reported in Table 4.3.

Table 4.3

Turbine Number 2 Exhaust - Sulfur Dioxide Emission Factor

Sulfur Dioxide Emission Factor, lbs/therm	0.0001*	
--	---------	--

^{*} Based on standard emission factor used by gas supplier (PG&E).

OUALITY ASSURANCE/QUALITY CONTROL

Source tests are performed to determine the types and amounts of pollutants emitted by a source. Information from this source test program may be used for obtaining permits, evaluating control equipment performance, updating emission inventories, and determining compliance with present and future emission regulations. For these purposes, reliable data are required. PES provides this reliability by using the following work practices:

USE OF STANDARD TEST PROCEDURES

CARB Methods 1 and 2 were utilized to measure flow rates. EPA Method 201A was used to determine the PM10 emission rate, and CARB Method 100 was used to determine the continuous emission rates for the gases. A procedure must be thoroughly studied under various conditions in order to be designated as a state or federal Method. Results of many executions of the procedure are compared to demonstrate accuracy and repeatability before adoption of the procedure as a source testing method.

USE OF TRAINED TEST PERSONNEL

Because of the complexity of typical source testing methods, testers should be trained and experienced with the test procedures in order to assure reliable results. PES personnel have had professional training and routinely conduct source tests.

KNOWLEDGE OF SOURCE'S OPERATION

The source testing team should have sufficient knowledge of the process to be tested in order to properly document the process parameters during the tests. Without documentation of the process parameters used, results are much less meaningful. PES has previously tested boilers and combustion sources and is familiar with the processes and equipment.

EQUIPMENT MAINTENANCE AND CALIBRATION

Use of properly maintained and calibrated test equipment is essential for minimizing systematic errors in results. All sampling devices were constructed,

maintained, and calibrated as suggested in EPA documents APTD-0576, and APTD-0581 (These are commonly accepted construction and maintenance manuals for source testing equipment). The dry gas meters were calibrated with a transfer gas meter with NBS traceability. These calibrations are included in Appendix D along with those for the nozzles, thermocouples, digital potentiometers, and Pitot tubes.

Quality control procedures used for continuous monitoring included the use of non-reactive 316 stainless steel or Teflon tubing and fittings throughout the system. A refrigeration unit was used with the pump down stream of the conditioned sample gas. All instrumentation was continuously monitored and checked between load conditions to insure data reliability during all sample runs. Bias checks were made with a calibration gas blend to confirm they met the tolerances specified in CARB Method 100.

All calibration gases were ± 1 per cent accuracy and provided by Scott Specialty Gases in San Bernardino, California. Copies of the calibration gas certifications are provided in Appendix D.

THOROUGH RECORD KEEPING

All data relating to the operation of the sampling train must be immediately recorded to ensure that it is not lost or misinterpreted. PES accomplishes this thorough record keeping by use of the field data sheets shown in Appendix B. The PES test team is familiar with these sheets and the information required to complete them. Any unusual occurrences in the process operation, unusual test instrument readings, or any other items that could affect the test results was also noted.

USE OF THOROUGHLY CLEANED GLASSWARE

All glassware and probe lines were cleaned prior to the tests with hot tap water and then with 40 per cent nitric acid solution. The trains were then cleaned with 0.1 Normal sodium hydroxide solution, laboratory grade distilled water, air dried, and sealed until the tests.

USE OF STANDARDIZED DATA REDUCTION TECHNIQUES

Data reduction was accomplished by the use of step by step calculation sheets. The calculations were systematic and easy to follow. All calculations for the source test are included in Appendix B.

SUBMISSION OF BLANK SAMPLES

Filter and reagent samples from an unused but charged PM10 sampling train carried to the field was analyzed with the other samples to detect any possible contamination of sampling media or problems with lab analyses. No corrections were made to the measured concentrations of the collected samples, but the blank train results were reported on the calculation sheets.



APPENDIX A EQUIPMENT PERMITS

— PACIFIC ENVIRONMENTAL SERVICES, INC. —



BAY AREA AIR QUALITY MANAGEMENT DISTRICT

939 ELLIS STREET SAN FRANCISCO, CALIFORNIA 94109 (415) 771-6000

Plant# 232

Page.

PERMIT

Expires: **JUL 1, 1993**

This document does not permit the holder to violate any District regulation or other town

Gerald Reid Onizuka Air Force Base P O Box 3430 Sunnyvale, CA 94088 ORIGINAL SENT TO: Onizuka Air Force Base 1080 Lockheed Way Sunnyvale, CA 94088

s#	DESCRIPTION	[Schedule]	PAID
1	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 1 Emissions at: Pl Stack	1 [B]	150
2	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 2 Emissions at: P2 Stack	1 [B]	150
3	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 3 Emissions at: P3 Stack	1 [B]	150
4	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 4 Emissions at: P4 Stack	1 [B]	150
5	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 5 Emissions at: P5 Stack	1 [B]	150
6	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 6 Emissions at: P6 Stack	[B]	150
7	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 7 Emissions at: P7 Stack	· [B]	150
8	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 8 Emissions at: P8 Stack	(B)	150
9	Turbine - Cogeneration, 12750K BTU/hr max, Multifue GAS TURBINE ENGINE WITH WASTE HEAT BOILER NO. 9 Emissions at: P9 Stack	[B]	150



APPENDIX B FIELD DATA AND CALCULATION SHEETS

PACIFIC ENVIRONMENTAL SERVICES, INC. —

TRAVERSE POINT LOCATION FOR CIRCULAR DUCTS

PLANT	ONIZUKA	AFB
DATE	12-2-	- 93
SAMPLING LOCATION	TURBI	NE 4
INSIDE OF FAR WALL TO OUTSIDE OF NIPPLE. (0	ISTANCE A)	25"
INSIDE OF NEAR WALL TO		l''
OUTSIDE OF NIPPLE. (DISTANCE A -	DISTANCE B)	24"
NEAREST UPSTREAM DIST	STURBANCE	2+0
CALCULATOR		24D

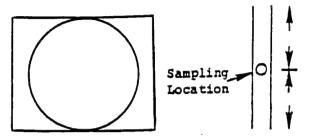
SCHEMATIC OF SAMPLING LOCATION

TRAVERSE POINT FRACTION NUMBER OF STACK I.D.		PRODUCT OF COLUMNS 2 AND 3 STACK I.D. (TO NEAREST 1/8 INCH)		TRAVERSE POINT LOCATION FROM OUTSIDE OF NIPPLE (SUM OF COLUMNS 4 & 5)	
0.067	24	1.6	·		
	1	60			
0.75	¥				
0.93		22.3			
	0.067 0.25	0.067 24 0.25 J	### COLUMNS 2 AND 3 COLUMNS 2 AND 3 CTO NEAREST 1/8 INCH	### COLUMNS 2 AND 3 OF STACK I.D. STACK I.D. (TO NEAREST 1/8 INCH) #### O. 25 O. 25 O. 75 18.0	

- PACIFIC ENVIRONMENTAL SERVICES, INC. -

PRELIMINARY VELOCITY TRAVERSE

Plant:	ONIZUKA AFB						
Date:	12-2-93						
Location:	TIRBINE #2						
Stack I.D.:	2+"						
	Pressure, in. Hg: 30.3						
Stack Gauge Pressure, in. H2O:							
Operators:	BROWN / LEARNEY						
	I.D. Number: Si7A						
Temperature	Readout I.D.: Box 3A						
Pitot Tube	Leak Check: OK						



Schematic of Traverse Point Layout

Traverse	Velocity	Stack	Cyclonic	Traverse	Velocity	Stack	Cyclonic
Point	Head (Aps)		Flow Check	Point	Head (Apg)	Temp.	Flow Check
Number	in. H ₂ O	(T.), °F	o from Null	Number	in. H ₂ O	(Tg), °F	° from Null
1 11	2.7	1:71				1	
1-4	2.7	673					
						[

						1	
	-						
						1	
						 	
					_	t	
Average				Average			
			- PACIFIC ENVIRON			<u> </u>	

LINER	\$2 × 0	लात.	18.00 18.00 14.70 14.70	6.80 E. 80 E	
- CLASS LINER	10 11 11 11 11 11 11 11 11 11 11 11 11 1	Im- pinger Temp.	1252	43 43 447 474 474 474 474 474 474 474 47	
55 - 5FT.	3. S.	Sample Box Temp. Filter Temp. °F	1 (1 1	1 , 3 1 1	
		Pump Vacuum In. fig	1444	1447 1447	
Length and Type	r Number	Dry Gas Meter Temp. Inlet Outlet 'Tmin' *F (Tmout) *F	527 6-1	69 69 71	
Probe L	Hitot Tube I Nozzie I.D. P Assumed Hoise Assumed Hoise Heter Box Nur Heter Gamma	Dry Gas H Inlet (Tmin) *F	59 71 73 73	27. 28. 38. 36. 1 _m T	
		Stack Temp. (Tg)	450	438 442 455 455 - - [4=60	
FIELD DATA	2 'R VIEW VIEW LIC OF OINE LAYOUT Local Control Control Local Control Loca	Pres. ntial . H20 Actual	22.0	2.	irks
FIEL	TOP VIEW Schematic of Traverse Point Layout Traverse Point Layout	Orifice Pres. Differential (AH) in. H20 Desired Actual	0.44	2.2 2.2 2.2 2.2 - -	2.45
	1.	Velocity Head (AP _B) in: H ₂ O	7.44	2.8 2.2 2.2 2.2 	502463 501641
AFB	47 - 30 6-1 - 21 6	Gas Heter Reading (Vm) ft	423,973 435.0 447.0 457.1 468.4	468.4 492.5 492.5 503.1 513.852	948121
CNIZUEA	Check Data	/Clock / Time /(24-hour clock)	07:19 07:49 08:03 08:18	08:24 08:39 09:54 09:08	00)× /
N.S	g Location Type ber Temperature Tremperature Tremperature Fressure Pressure (Pg) Number(g) Leak Rate = Pitot Leak C	Sampling Time, (min)/	15 15 30 28.8 58.8	58.3 74.1 834.1 103.0 116.1	METER 07:13 39 40
Plant	Sampling Location 1025 Sample Type Run Runder Operator Ambient Temperature Barometric Pressure Static Pressure Filter Number(s) Pretest Leak Rate = 0.06 Pretest Pitot Leak Check Pretest Orsat Leak Check Read and Record all Data	Traverse Point Number	ス (1) 20 20 20 20 20 20 20 20 20 20 20 20 20	(i) 23 FINI	Ú45
		PACIFIC ENVIRO	DNMENTAL SERVIC	es. INC.	



SAMPLE RETRIEVAL DATA

Plant:		DNIZU	IKA AFE	3			
Date:		12	12-3-93				
Sampling Location		TU	RBINE #2	STAC	K		
Sampling Type (M	ethod):		PM-10				
Run Number: 38			MAD 0	AFB-	(
Sample Box Number	r:		3B	A () . (c) . (
Clean-up Man: Job Number:			BROWN KE	arney			
Commontes			F028				
Filter				•			
Filter Number: Description of F	ilter: <u>CUE</u>	₹N					
Moisture	#		#2-		* 3		
<pre>Impingers: Final Volume:</pre>	1570	 7	102.0	mL	16.0	T	
Initial Volume:	100.0	mL mL	100.0	mL _	0.0	mL mL	
Net Volume:	57.0	mL	2.0	mL _	16.0	mL	
Total H ₂ O:	57	mL	59	_mL _	75	mL	
Silica Gel							
Final Volume:	700.9	g _		g _		g	
Initial Volume:	681-9	g		g _		g	
Net Volume:	19.0	g _	······································	g _		g	
Total Moisture:	94	g _		g _		—а	
Description of In	pinger catch	:	CLEAR				
			1				

PACIFIC ENVIRONMENTAL SERVICES, INC.

Date: 12-3-93

Source/Sample Number: OAFB-1

- 1. $Vm(std) = (17.64)(Vm)(Y) \left[\frac{P_{bar} + (\Delta H/13.6)}{Tm} \right]$ $Vm(std) = (17.64)(89.879)(0.99) \left[\frac{(30.2) + (2.15/13.6)}{(52.9)} \right]$ $Vm(std) = 90.1 \quad dscf.$
- 2. Volume water vapor collected (standard conditions).

V(10) = 94 condensate from impingers and selica gel.

$$Vw(std) = (0.04707) V(10) = (0.04707) (9)$$

 $Vw(std) = \underbrace{Y.42}_{scf.}$

3. Percent moisture, by volume.

$$Bw_s = \frac{Vw(std)}{Vw(std) + Vm(std)} = \frac{(4.42)}{(4.42) + (90.1)} = \frac{0.047}{}$$

 $BW_{S} = 4.71.$

4. Molecular weight, stack gas.

Dry molecular weight.

$$Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$$

$$Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80)$$

 $Md = 29.1 \quad 1b/1b-mole.$

$$Ms = Md + Bw_s (18 - Md) = (29.1) + (0.647)(18 - 29.1)$$

Ms = 28.58 lb/lb-mole.

Date: 12-3-93

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(V\Delta P) \left[avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(0.84)(1.59) \left[\frac{(910)}{(30.1)(28.58)} \right]$$

Vs(avg) = 117.4 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(117.4)(3.14)$$

Qs = 22118 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

$$Q(std) = (17.64)(Qs)(1-Bw_s) \frac{(Ps)}{(Ts)}$$

$$Q(std) = (17.64)(22/18)(1 - 0.847) \begin{bmatrix} 30.1 \\ 910 \end{bmatrix}$$

 $Q(std) = 12299 \quad dscfm.$

8. Isokinetic variation.

$$2I = (K) \left[\frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$

$$z_{I} = (0.0945) \left[\frac{()()}{()()()(1-)} \right]$$

$$xI = \sqrt{A}x$$



Date: 12-3-43

9. Viscosity of stack gas:

$$\mu_s = 152.418 + 0.2552 t_s + 3.2355 \times 10^{-5} (t_s)^2 +$$

0.53147 (%O₂) - 74.143 Bw_s

$$\mu_s = 152.418 + 0.2552(450) + 3.2355x10^{-5} (450)^2 +$$

0.53147(17.5) - 74.143(0.047)

 $\mu_s = \frac{279.0}{}$ micropoise.

10. Cyclone flow rate:

$$Q_s = 0.002837 \ \mu_s \ \left| \frac{t_s + 460}{M_w P_s} \right|^{0.2949}$$

$$Q_{s} = 0.002837 (279.6) \left| \frac{(450) + 460}{(23.58)(30.1)} \right|^{0.2949}$$

 $Q_{n} = 0.81$ ft³/min

11. Orifice pressure head for cyclone flow rate:

$$\Delta H = \left(\frac{Q_s (1-Bw_s) P_s}{t_s + 460} \right)^2 \left(\frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$$

$$\Delta H = \left(\frac{(0.0)(1-0.047)(30.1)}{(.450) + 460} \right)^{2} \left(\frac{(69 + 460)(29.7)(1.083)(7.976)}{(30.2)} \right)$$

 $\Delta H = 0.69$ in. H_2O



Date: 12-3-93

Source/Sample Number: <u>OAFB-I</u>

Stack viscosity, μ_s , micropoise = $\frac{279.6}{30.1}$ Absolute stack pressure, P_s , in. Hg = $\frac{30.1}{30.1}$ Average stack temperature, t_s , $F = \frac{450}{69}$ Method 201A pitot coefficient, $C_s = \frac{69}{0.84}$ Cyclone flow rate, ft³/min, $Q_s = \frac{0.81}{0.14}$ Method 2 pitot coefficient, $C_s = \frac{0.81}{0.14}$ Molecular weight of stack gas, wet basis, $M_s = \frac{28.58}{0.150}$ Nozzle diameter, D_s , in. = $\frac{0.150}{0.150}$

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = 110.0 \, \text{ft/sec}$$

Maximum and minimum velocities:

Calculate R_{min}.

If R_{min} is less than 0.5, or if an imaginary number occurs when calculating R_{min} , use Equation 1 to calculate v_{min} . Otherwise, use Equation 2.

Eq. 1
$$v_{min} = v_n (0.5) = ____ ft/sec$$

 $v_{min} = v_n R_{min} = _{82-5} ft/sec$

Calculate R_{max} .

$$R_{\text{max}} = 0.4457 + \sqrt{0.5690 + \frac{0.2603 (\sqrt{Q_s}) \mu_s}{v_n^{1.5}}} = 1.24$$



Plant: ONIZUKA AFB

Date: 12-3-9.3

Source/Sample Number: OAFB-1

If R_{max} is greater than 1.5, use Equation 3 to calculate v_{max} . Otherwise,

Eq. 3
$$v_{max} = v_n (1.5) = ____ ft/sec$$

Eq. 4 $v_{max} = v_n R_{max} = _136.4$ ft/sec

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = \frac{1.25}{10.00} in. H_20$$

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = \frac{3.41}{\text{in. H}_20}$$

Calculate the actual D_{50} of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left(\frac{t_s + 460}{M_w P_s} \right)^{0.2091} x \left(\frac{\mu_s}{Q_s} \right)^{0.7091}$$

where, $\beta_1 = 0.15625$

$$D_{50} = (0.15625) \left(\frac{910}{(2858)(30.1)} \right)^{0.2091} \times \left(\frac{279.6}{0.81} \right)^{0.7091}$$

$$D_{50} = 9.97 \mu m$$



CLIENT: ONIZUKA AFB

Project No. <u>F028</u>

PM 10 TEST *1

PARTICULATE CALCULATIONS

Sampling Loc	ation: _	TURBINE	#2 EXHAUST	Test Date: /	12-03-93				
Sample Numbe	Sample Number: <u>OAF8-/</u> Sample Volume: <u>90./</u> DSCF								
Stack Flow R	Stack Flow Rate: 12,300 DSCFM								
Particulate	Catch:	(grams	PM10	- PM10					
Filter:	Initial	Weight Weight Weight	0.2084 0.2083 0.000						
Impinger:	Initial	Weight Weight Weight	28.6871 28.6878 -0.0007	29.0835 29.0832 0.0003					
Extract:	Initial	Weight Weight Weight							
Total:			<0.0010	40.0010					

Particulate Concentration:

 $\frac{(40.00) \text{ grams x } 15.43 \text{ grains/gram}}{(90.) \text{ DSCF}} = \frac{60.0002}{\text{grains/DSCF}}$

Particulate Emissions:

 $\frac{(20.000)}{9000} \frac{\text{gr/DSCF} \times (/2/300)}{1000} \frac{\text{DSCFM} \times 60 \text{ min/hr}}{1000} = \frac{20.02}{1000} \frac{1}{1000} \frac{1}{100$

Rule 404 Limitation @ _____ SCFM = ____ grains/DSCF

Rule 405 Limitation @ NA = 1bs/hr = 1bs/hr

— PACIFIC ENVIRONMENTAL SERVICES, INC. —

A 50 0 0 in. lig	MIN 0.21 0.21 72.81 72.81	5.7.7.7.7.7.7.7.7.7.7.7.7.7.7.7.7.7.7.7	
1 In. 11g	Im- Im- Temp. °F %	58	9 9 1
3.A 1.916 0.69 0.69 0.69 0.69 0.69	Sample Box Temp. Filter Temp. °F	1 1 1 1	
Type SS	> -1 1 1 1	20021	
Bth and Te I.D. No. 6 olsture, Number Number Apple Leak Rapitot L. Orsat L.	Dry Gas Heter Temp. Inlet Outlet Tmin *F (Tmout) *F 74 70 83 74 70 93 93 74	\$8 88 87 87 87 87 87	87
Probe Length Pitot Tube I. Nozzle I.D. n Assumed Hoist Heter Alle C Factor Heter Gamma Heter Gamma Heter Gamma Heter Gamma Heter Gamma Fost Test Le Fost Test De Fost D	Inlet (Tmin) °F	95 100 48 96	<u> </u>
		464	15.451
Schematic of Erse Point Layout Layout Layout	Pres. ntla1 1. 1120 Actual 2.2 2.2 2.3	4.00 C.	2.3 Hrs
(SEE TEST 1) Schematic of Traverse Point Layout Traverse Point Layout	Orifice Pres. Differential (All) in. 1120 Desired Actual 2.2 2.2 2.2 2.2 2.3 2.3 2.3 2.3	4.2.4.4.4.4.4.4.4.4.4.4.4.4.4.4.4.4.4.4	2H 2
	Velocity Head (APB) In: H20 2.7 2.7 2.7 2.7 2.7 2.7	य प्राप्त प्र	102641 302641 32495 264
2000 2000 2000 2000 2000 2000 2000 200	Gas Meter Reading (Vm.) ft 513.919 524.3 536.4 541.7 559.8	558.8 571.6 534.3 595.7 607.011	9 4 8 363 9 4 8 107 344
Check Check Check	/Clock / Time /(24-hour clock) / 10:30 / 10:45 / 10:45 / 1:14		00)
ing Location The Type unber tor nt Temperature etric Pressure c Pressure (Pg) ir Number(s) ir Number(s) st Leak Rate = 6.0 sst Oreat Leak Che sst Oreat Leak Che and Record all Da	Sampling Time, (min) / 0 15.0 30.0 43.5	\$7.0 7 6 .3 8 9 .0 10[.1	METER 04:40 13:04
Plant CNN ZL Bate Sampling Location Sample Type Run Rumber Operator Ambient Temperature Barometric Pressure (Ps Filter Number(s) Pretest Leak Rate Pretest Ditot Leak Pretest Orsat Leak Read and Record all	Traverse Point Numbor L H 2 2 (1) 2 FINI	R (1)	ëA5
	PACIFIC ENVIRCINMENTAL SERVICE	S, INC.	



SAMPLE RETRIEVAL DATA

Plant:	ONIZUKA /					
Date:	12-3-9	12-3-93				
Sampling Location:	TURBINE	#2 STACK				
Sampling Type (Method): _	PM-	10				
Run Number:	OAF	3-2				
Sample Box Number:	3F	3				
Clean-up Man:	BROWN	KEARNEY				
Job Number:		8				
Comments:						
	•					
	,					
Filter	***************************************	•				
	•					
		•				
Filter Number:						
Description of Filter:	CLEAN					
•						
Moisture						
*	(#2	* 3			
		. •				
Impingers:						
Final Volume: 156	.0 mL	108 mL	3.0 mL			
Initial Volume: 100	.O mL	100.0 mL	0.0 mL			
Net Volume: 56	mL	8 mL —				
Total H ₂ O: 50		64mL _	67 mL			
10ta1 n ₂ 0.						
	1					
Silica Gel	Ý.					
Final Volume: 68	<u>2.3</u> 'g	g _	g			
Initial Volume: 65	9.7 g	g	g			
	2.6 g	g	g			
Total Moisture: \$9	.6 g	g _	g			
	ľ					
Description of Inpinger c	atch:CLE	tr				
	1					
	:					
	4					

PACIFIC ENVIRON MENTAL SERVICES, INC.



Date: 12-3-93

Source/Sample Number: <u>OAFB-2</u>

- 1. $Vm(std) = (17.64)(Vm)(Y) \left[\frac{P_{bar} + (\Delta H/13.6)}{Tm} \right]$ $Vm(std) = (17.64)(93.692)(6.99) \left[\frac{(3c2) + (2.3)(13.6)}{(547)} \right]$ $Vm(std) = 90.3 \quad dscf.$
- 2. Volume water vapor collected (standard conditions).

V(10) = 89.6 condensate from impingers and selica gel.

Vw(std) = (0.04707) V(10) = (0.04707)(89.6)

Vw(std) = 4.22 scf.

3. Percent moisture, by volume.

 $BW_S = \frac{VW(std)}{VW(std) + Vm(std)} = \frac{(4.22)}{(4.22) + (90.3)} = \frac{0.045}{}$

 $Bw_s = 4.5 i$.

4. Molecular weight, stack gas.

Dry molecular weight.

 $Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$

Md = 0.440 (2.5) + 0.320(7.5) + 0.280(80)

Md = 29.1 lb/lb-mole.

 $Ms = Md + Bw_s (18 - Md) = (29.1) + (0.045)(18 - 29.1)$

Ms = 78.6 1b/1b-mole.



Date: 12-3-93

Source/Sample Number: <u>OAFB-2</u>

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(V\Delta P) \left[avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(0.87)(1.58) \sqrt{\frac{917}{(30.1)(28.6)}}$$

Vs(avg) = 117.1 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(117.1)(3.1 \neq)$$

Qs = 22062 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

Q(std) =
$$(17.64)(Qs)(1-Bw_s)[(Ps)]$$

$$Q(std) = (17.64)(22062)(1 - 0.845) \begin{bmatrix} 30.1 \\ 917 \end{bmatrix}$$

Q(std) = 12200 dscfm.

8. Isokinetic variation.

$$2I = (K) \left[\frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$

$$z_{I} = (0.0945) \left[\frac{()()}{()()()(1-)} \right]$$



Date: 12-3-93

Source/Sample Number: OAFB-2

9. Viscosity of stack gas:

$$\mu_s = 152.418 + 0.2552 t_s + 3.2355x10^{-5} (t_s)^2 + 0.53147 (%O_2) - 74.143 Bw_s$$

$$\mu_s = 152.418 + 0.2552(457) + 3.2355 \times 10^{-5} (457)^2 + 0.53147(17.5) - 74.143(0.045)$$

 $\mu_* = \frac{28i.8}{}$ micropoise.

10. Cyclone flow rate:

$$Q_s = 0.002837 \ \mu_s \ \left| \frac{t_s + 460}{M_w P_s} \right|^{0.2949}$$

$$Q_{s} = 0.002837(281.8) \left[\frac{(375) + 460}{(28.6)(30.1)} \right]^{0.2949}$$

 $Q_a = 0.81$ ft³/min

11. Orifice pressure head for cyclone flow rate:

$$\Delta H = \left(\frac{Q_s (1-Bw_s) P_s}{t_s + 460} \right)^2 \left(\frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$$

$$\Delta H = \left| \frac{(0.81)(1-0.045)(30.1)}{(457) + 460} \right|^2 \left| \frac{(87 + 460)(29.1)(1.083)(1.916)}{(30.2)} \right|$$

 $\Delta H = \delta \cdot 71$ in. H_2O



Date: 12-3-93

Source/Sample Number: OAFB-2

Stack viscosity, μ_s , micropoise = $\frac{281.8}{30.i}$ Absolute stack pressure, P_s , in. Hg = $\frac{30.i}{457}$ Meter temperature, t_s , *F = $\frac{457}{87}$ Method 201A pitot coefficient, C_s = $\frac{0.84}{100}$ Cyclone flow rate, ft /min, Q_s = $\frac{0.81}{100}$ Method 2 pitot coefficient, C_s = $\frac{0.84}{100}$ Molecular weight of stack gas, wet basis, M_s = $\frac{0.150}{100}$

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = \frac{110.0}{\text{ft/sec}}$$

Maximum and minimum velocities:

Calculate R_{min}.

If R_{\min} is less than 0.5, or if an imaginary number occurs when calculating R_{\min} , use Equation 1 to calculate v_{\min} . Otherwise, use Equation 2.

Eq. 1
$$v_{min} = v_n (0.5) =$$
_____ ft/sec
Eq. 2 $v_{min} = v_n R_{min} =$ _82-5 ft/sec
Calculate R_{max} .

$$R_{\text{max}} = 0.4457 + \sqrt{0.5690 + \frac{0.2603 (\sqrt{Q_s}) \mu_s}{v_n^{1.5}}} = 1.24$$



Date: 12-3-93

Source/Sample Number: <u>6AF6-2</u>

If R_{max} is greater than 1.5, use Equation 3 to calculate $v_{\text{max}}.$ Otherwise, use Equation 4.

Eq. 3
$$v_{max} = v_n (1.5) = ____ ft/sec$$

Eq. 4 $v_{max} = v_n R_{max} = ___ 136.4$ ft/sec

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = 1.27$$
 in. H₂0

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = 3.39 \text{ in. } H_20$$

Calculate the actual D_{50} of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left(\frac{t_s + 460}{M_w P_s} \right)^{0.2091} x \left(\frac{\mu_s}{Q_s} \right)^{0.7091}$$

where, $\beta_1 = 0.15625$

$$D_{50} = (0.15625) \left(\frac{917}{(28 \cdot 6)(36)} \right)^{0.2091} \times \left(\frac{281.8}{0.81} \right)^{0.7091}$$

$$D_{50} = 16.5 \% \mu m$$



CLIENT: ONIZUKA AFB
Project No. F028

	PARTICUI	ATE CALCULATI	ONS PMIO	HEST #2	
Sampling Location: _	TURBIN	E #2 EXHAUST	Test Date: _	12-03-94	
Sample Number:	48-2	Sample Volu	me: 90.3	DSCF	
Stack Flow Rate:	12,200	DSCFM	·		
Particulate Catch:	(gram	s) PM10	> PMID		
Initial	Weight	0.2066 0.2072 -0.0006			
Initial	Weight	28.6117 28.6117 0.0000	28.3456 28.3443 0.0013		
Initial	Weight Weight Weight				
Total:		20.0010	0.0013		
Particulate Concentr		<u>grains/gram</u> SCF	= <0.0002	grains/DSCF	
Particulate Emissions: $(\angle 0,0002)$ gr/DSCF x $(12,200)$ DSCFM x 60 min/hr = $\angle 0.02$ lbs/hr 7000 grains/lb					
Rule 404 Limitation	e <u>NA</u>	SCFM =	grains/	DSCF	
Rule 405 Limitation	e <u>NA</u>	lbs/hr =	lbs/h	ır	

PACIFIC ENVIRONMENTAL SERVICES, INC. --



SAMPLE RETRIEVAL DATA

Plant:	ONIZI				
Date:	(2-3-93				
Sampling Location:		TURBINE !		CK	
Sampling Type (Method):		PM-11	0		
Run Number: Sample Box Number:		OAFB	- 3		
		3B	KEARA	1611	
Clean-up Man:		BROWN FO2	2 HEARA	UEY	
Job Number:		F0Z	δ		
Comments:					
			-		
Filter					
Filter Number:					
Description of Filter:					
Moisture					
Impingers:	,				
Final Volume:	54. b	_mL	106.0	mL	#3.0 mL
	00.0	_mL	100.0	mL	0.0 mL
	54	_mL	6.0	mL	3.0 mL
Total H ₂ O:	54	_mL	60	mL	63 mL
Silica Gel					
Final Volume:	715.2	a		_g	g
	690.1	_g		_g	g
Net Volume:	25.1	_g		_g	g
Total Moisture:	88.1	_g		g	g
Description of Inpinger	catch:				

PACIFIC ENVIRONMENTAL SERVICES, INC.

Date: 12-3-93

Source/Sample Number: OAFB-3

- 1. $Vm(std) = (17.64)(Vm)(Y) \left[\frac{P_{bar} + (\Delta H/13.6)}{Tm} \right]$ $Vm(std) = (17.64)(91.077)(0.99) \left[\frac{(30.2) + (2.2)(13.6)}{(592)} \right]$ $Vm(std) = 89-1 \quad dscf.$
- 2. Volume water vapor collected (standard conditions).

V(10) = 88.1 condensate from impingers and selica gel.

$$Vw(std) = (0.04707) V(10) = (0.04707)(884)$$

Vw(std) = 4-15 scf.

3. Percent moisture, by volume.

$$Bw_{s} = \frac{Vw(std)}{Vw(std) + Vm(std)} = \frac{(4.15)}{(4.15) + (89.1)} = \frac{0.045}{}$$

Bws = 4.5%.

4. Molecular weight, stack gas.

Dry molecular weight.

$$Md = 0.440(\% CO_2) + 0.320(\% O_2) + 0.280(\% N_2 + \% CO)$$

$$Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80)$$

Md = 291 lb/lb-mole.

$$Ms = Md + Bw_s (18 - Md) = (29.1) + (0.045)(18 - 29.1)$$

Ms = 28.0 lb/lb-mole.

Date: 12-3-93

Source/Sample Number: <u>OAFB-3</u>

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(\sqrt{\Delta P}) \left[avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(0.84)(1.56) \sqrt{\frac{920}{(30.1)(28.6)}}$$

Vs(avg) = 115.8 ft/sec.

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

$$Qs = (60)(Vs)(A) = (60)(115.8)(3.14)$$

Qs = 21817 acfm.

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

Q(std) =
$$(17.64)(Qs)(1-Bw_s)[(Ps)]$$

$$Q(std) = (17.64)(21817)(1 - 0.045) [30.1]$$

Q(std) = 1202-5 dscfm.

8. Isokinetic variation.

$$2I = (K) \left[\frac{(Ts)(Vm(std))}{(Ps)(Vs)(An)(\theta)(1 - Bw_s)} \right]$$

$$z_{I} = (0.0945) \left[\frac{()()}{()()()(1-)} \right]$$



Date: 12-3-93

Source/Sample Number: <u>OAFB-3</u>

9. Viscosity of stack gas:

$$\mu_s = 152.418 + 0.2552 t_s + 3.2355 \times 10^{-5} (t_s)^2 +$$

0.53147 (%O₂) - 74.143 Bw_s

 $\mu_s = 152.418 + 0.2552(460) + 3.2355 \times 10^{-5} (460)^2 +$

0.53147(17.5) - 74.143(0.045)

 $\mu_{\text{m}} = 282 - 6$ micropoise.

10. Cyclone flow rate:

$$Q_s = 0.002837 \ \mu_s \ \left(\frac{t_s + 460}{M_c P_s}\right)^{0.2949}$$

$$Q_{a} = 0.002837 (282.6) \left| \frac{(460) + 460}{(78.6)(30.1)} \right|^{0.2949}$$

 $Q_{\bullet} = 0.82 \text{ ft}^3/\text{min}$

11. Orifice pressure head for cyclone flow rate:

$$\Delta H = \left(\frac{Q_s (1 - Bw_s) P_s}{t_a + 460} \right)^2 \left(\frac{(t_m + 460) M_d (1.083) \Delta H@}{P_{bar}} \right)$$

$$\Delta H = \left(\frac{(0.82)(1-0.045)(30.1)}{(4400) + 460} \right)^{2} \left(\frac{(82 + 460)(29-1)(1.083)(1.916)}{(30.2)} \right)$$

 $\Delta H = 0.71$ in. H_2O



Date: 12-3-93

Source/Sample Number: <u>DAFB-3</u>

Stack viscosity,
$$\mu_s$$
, micropoise = 282.6

Absolute stack pressure, P_s , in. Hg = 30.1

Average stack temperature, t_s , $F = 460$

Meter temperature, t_s , $F = 82$

Method 201A pitot coefficient, $C_s = 0.84$

Cyclone flow rate, ft³/min, $Q_s = 0.84$

Method 2 pitot coefficient, $C_s = 0.84$

Molecular weight of stack gas, wet basis, $M_s = 28.6$

Nozzle diameter, D_s , in. = 0.150

Nozzle velocity

$$v_n = \frac{3.056 \, Q_s}{D_n^2} = \frac{11.4}{\text{ft/sec}}$$

Maximum and minimum velocities:

Calculate R_{min}.

If R_{min} is less than 0.5, or if an imaginary number occurs when calculating R_{min} , use Equation 1 to calculate v_{min} . Otherwise, use Equation 2.

Eq. 1
$$v_{min} = v_n (0.5) = ____ ft/sec$$

Eq. 2 $v_{min} = v_n R_{min} = _83.6 ft/sec$

Calculate R_{max} .



Date: 12-3-9.3

Source/Sample Number: <u>OAFB-3</u>

If R_{max} is greater than 1.5, use Equation 3 to calculate $v_{\text{max}}.$ Otherwise, use Equation 4.

Eq. 3
$$v_{max} = v_n (1.5) = ____ ft/sec$$

Eq. 4 $v_{max} = v_n R_{max} = _{138.1} ft/sec$

Maximum and minimum velocity head values:

$$\Delta p_{min} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{min})^2}{(t_s + 460) C_p^2} = 1.27 \text{ in. } H_20$$

$$\Delta p_{\text{max}} = 1.3686 \times 10^{-4} \frac{P_s M_w (v_{\text{max}})^2}{(t_s + 460) C_p^2} = 3.46 \text{ in. } H_20$$

Calculate the actual D_{50} of the cyclone for the given conditions as follows:

$$D_{50} = \beta_1 \left(\frac{\underline{t}_s + 460}{\underline{M}_w P_s} \right)^{0.2091} \times \left(\frac{\underline{\mu}_s}{Q_s} \right)^{0.7091}$$

where, $\beta_1 = 0.15625$

$$D_{50} = (0.15625) \left(\frac{920}{(28 \cdot 6)(30 \cdot i)} \right)^{0.2091} \times \left(\frac{282 \cdot 6}{0.32} \right)^{0.7091}$$

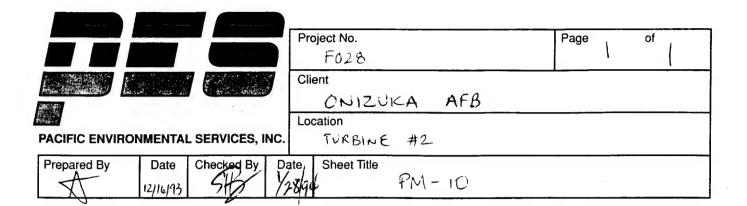
$$D_{50} = 9.98 \mu m$$



Project No. F028

			•			
			IONS PMIO			
Sampling Loca	ation: TURBINE	#2 EXHAUST	Test Date: _	12-03-94		
Sample Number	r:	Sample Volu	ume: 89.1	DSCF		
Stack Flow Ra	ate: 12,030	DSCFM				
Particulate (Catch: (gram	s) PMIO	> PMIO			
Filter:	Final Weight Initial Weight Net Weight	0.2056 0.2065 - 0.0009				
Impinger:	Final Weight Initial Weight Net Weight	28.6882	28.1727 28.1721 0.0006			
Extract:	Final Weight Initial Weight Net Weight					
Total:		20.0010	0,0006			
Particulate (Concentration:					
$\frac{(20.00/0) \text{ grams x 15.43 grains/gram}}{(89.1) \text{ DSCF}} = \frac{<0.0002}{\text{grains/DSCF}}$						
Particulate E	missions:					
(20,000) gr	/DSCF x (/2,030 7000 grain) DSCFM x 60 ns/lb	$min/hr = \frac{\angle 0.6}{2}$	02 lbs/hr		
Rule 404 Limi	tation @ NA	SCFM =	grains/I	SCF		
Rule 405 Limi	tation @ NA	lbs/hr =	lbs/hr	•		

-- PACIFIC ENVIRONMENTAL SERVICES, INC. -



BEAKER	POST WEIGHT &	PRE WEIGHT 9	NET g
1 A	29.0035	29.0832	0.0003
2A	28.3456	28.3443	0.0013
3A	28.1727	28.1721	0.0006
18	28.6871	28. 6£78	-0.0007
28	28-6117	28.6117	0. 0000
3B	28.6882	23. 5382	0.0000

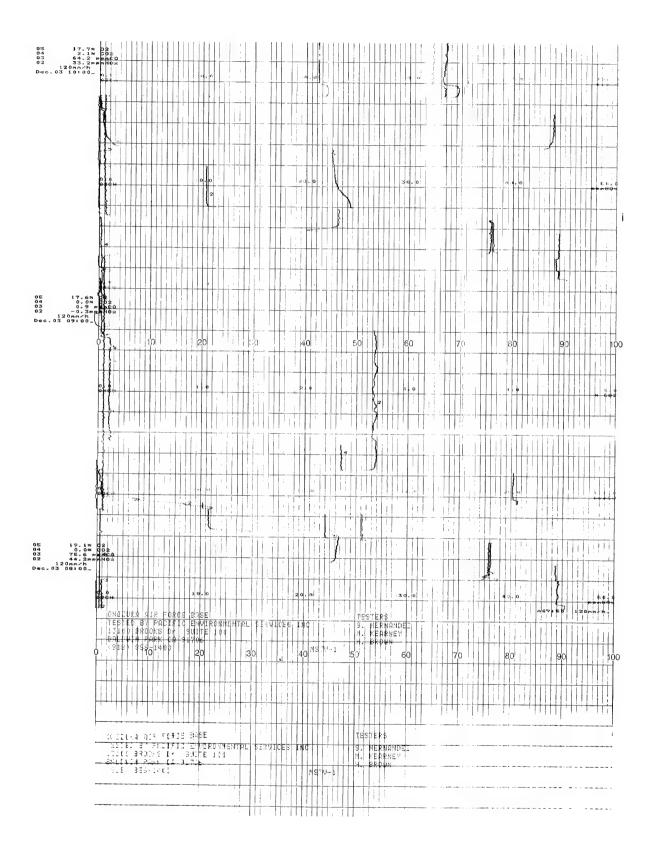
NOTE: NO SIGNIFICANT CATCHES

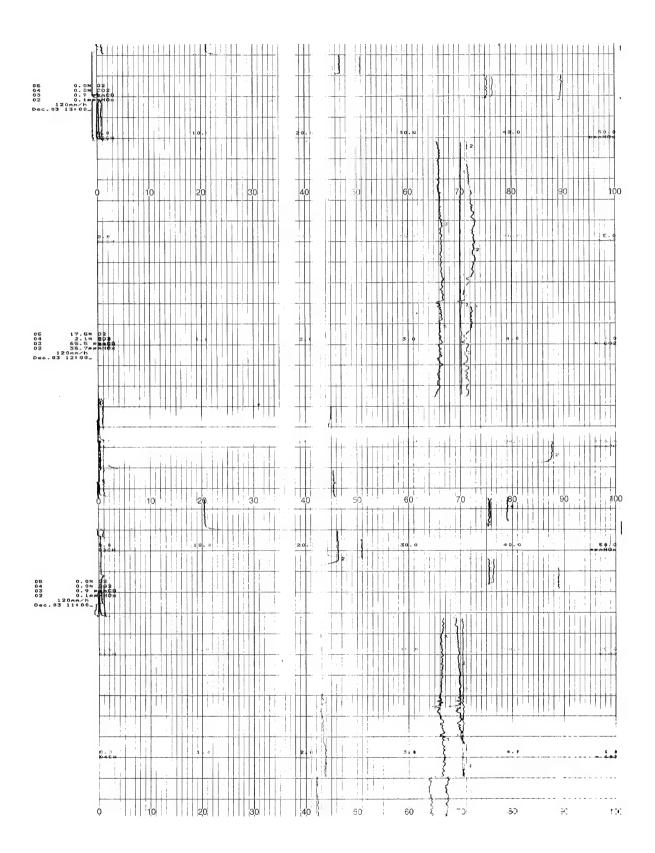
1.E. < 1 mg PM10 < 1 mg > PM10

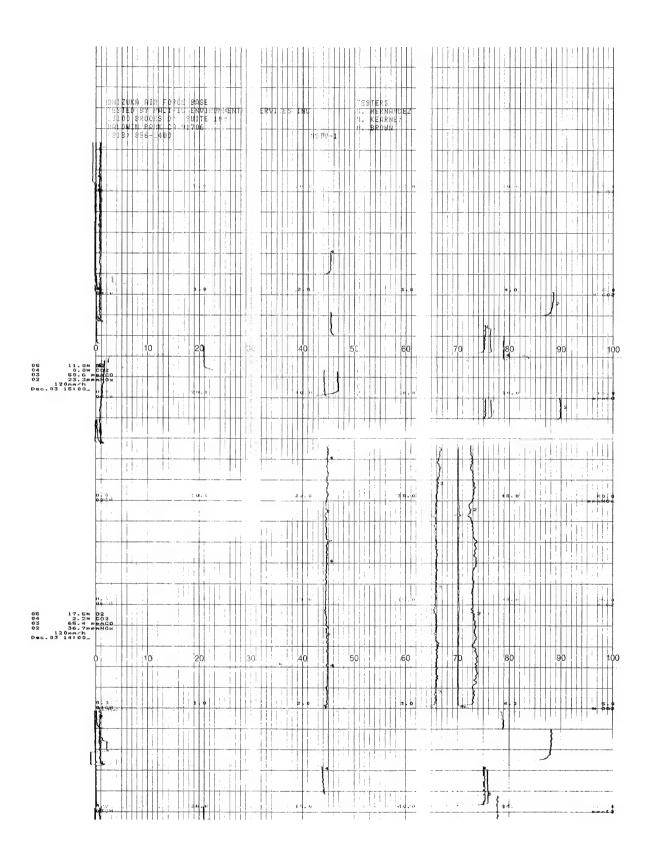
FILTERS :

1	0.2084	0.2083	0.0001
2	0 2066	0.2012	-0.0006
3	0.2056	0-2065	-0.0009

0% 0.0 04 0.0 02 0.0 02 12000 Dec. 02 17700	0 10	20	1.00			93
	O	20		50 50 E ### E #### E ######	 80	90 100 90 100







			Project No. Page 1 of 8	
			Client CNIZUKA, AEB	
PACIFIC ENVIRONMENTAL SERVICES, INC.			CONTINUE / CA	į
Prepared By	Date 12-ろ	Checked By Ы/Д	Date Sheet Title VAC & START 19.8"45 - 19.8" & GARB 2	(مر

. -		C >	COL	02	NOTES.
TIME	Nox	co			No.
	0.0	0,9	0.0	0.0	
	44.2	75.6	49,0	19,0	
	23.1	51.0	2. 3	11,0	N. SPAN 22-3
-	10,7				L- SPAN 10.4
08:23	24.8				CONVERTER CHEK (NO,)
08:53	26.5				On Convoctor CHOIL
n 0 \ n .		_	_		OL TRAVORSE R. B (4) SE
08:26				17.6	
				17.6	R (3) R (4) R (4) R (3) R (4) R (4) R (4) R (3)
				17.6	A- (1)
		-		17.6	R 5 (1)
				17.6	39/120 R 3 (3)
				17.6	3/ /
				17. 7	R 2 (2)
~ ~				17.6	/ R ((1)
- -	પ ર. છ	75.5	2.2	19.0	Bias (HIGH)
57827	eun 1				
09:56	34.7	لولد. ك	7.1	17.5	e R4 F/5 W2= 150
09:57	34. Z	64.6	21	17.5	@ C.250 VOC
01:58	34.0	65.2	2-1	17.6	•
09.59	3 3, 1	64.2	2.1	17.7	
€0:00	33. 4	64.2	2.1	17-7	
10.01	33,2	44.1	2.1	17.7	
10:02	33.3	64.2	2.1	17.7	
10:03	<i>3</i> 3, 4	64.4	2.1	17.7	
10.64	33,5	64.4	2.1	17.7	
101.05	33, H	64.1	2.1	17.7	

				Project No. Page of 2 8				
				Client ONIZUKA AFB				
PACIFIC ENVIRONMENTAL SERVICES, INC.				Sunny sue, LA				
Prepared By	Date	Checked By	Date	Sheet Title	N 1			

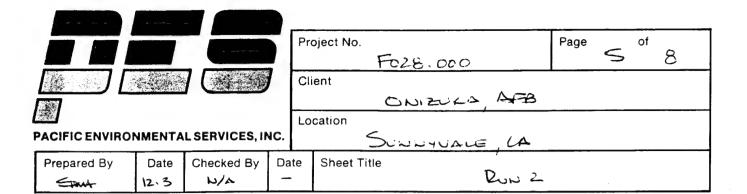
TIME	Nox	co	C02	02_	NOTES:
10:06	33.5	64.4	144	17.7	
FO:01	33.5	64.3	2.1	17.7	
10108	33,5	64.3	211	127	
10:09	33.5	64.3	2.1	12.7	
10:10	33.6	64.1	2.1	17,7	HADUSO to Q 3 SW
10:11	33.6	64.2	2.1	17.7	
10/12	33.6	64.2	2.1	17.7	
10:13	- 33. 7	44.3	2.1	12.7	
10:14	35.7	64.1	2.1	17.7	
10:15	33.6	63.8	2.0	17.7	(Zo niu)
10:16	33.5	63,9	2.1	17.7	
10:17	33.7	64.1	2.1	17.7	SCTUAL POINT CHANGE
10:18	34.6	65.2	2.1	17.6	CHANGES OBSERVED ON
10:19	35.2	66.4	2.1	17.6	STORP CHART.
10:20	35.3	46.7	2.1	17.6	
10:21	35.5	66.9	2.1	17,5	
10:22	35.4	66.6	2.1	17.5	
10:23	35.5	64.5	2.1	17.5	
10:24	35.5	44.5	2.1	17.5	
10: 25	35.5	46,5	2.1	17.5	(30 MIN)
10:26	3 5, 3	44.5	2-1	172	
10127	35.2	66.5	2.1	17.6	
10:28	55.1	Lele. Le	2.1	13.2	
10129	35.0	66.6	2.1	17.5	A CONTRACTOR OF THE PROPERTY O
10130	34,7	66.3	2.1	17.6	Commission of the Commission o
10:31	34.7	66.0	2.1	17.6	
10:32	35.0	66.0	2.1	176	
10:33	35.0	66.0	2.1	17.6	
10:34	34,6	65,6	2.1	15.0	
10.35	34.9	45.8	21	17.6	(40 mm)

				F	Project No. Page of 3 8	Page of	
				7	Client ONIZUKA APB	_	
•	PACIFIC ENVIRONMENTAL SERVICES, INC.				Suinguace, CA		
	Prepared By	Date \2、3	Checked By	Date	•		

TIME	Nov	<u>co</u>	CO2	02	Notes :
10136	34.1	45.8	2.1	17.0	
10:31	349	45.9	2.(17.6	
10:38	34.8	65.9	2.1	17.6	4 4000 40 Pc 3
10:39	34.8	65.8	2.1	17.6	
10:40	34.8	66.0	2.1	17.6	
10:41	34.9	66.2	2.1	17.6	
10:42	34.8	662	2.1	17.6	
10:43	34.8	66.4	2.1	17.6	
10:44	34.9	66.3	2.1	17.5	
10:45	34.8	46.3	2.1	17.6	(St MIN)
10:46	34.8	66.4	2.1	17.5	
10:47	ろ り、子	66.2	2.1	34.5	
10:48	34,7	66.5	2.1	17.5	
10:49	34.8	46.2	2.1	17.6	
10:50	34,7	662	2.1	17.0	
10:57	34.7	46.0	2.1	17.6	,
10:52	34.7	أدلورا	21	13.6	(x co =
10:53	34, 6	ا، ناب	2.1	17.6	× 602=
10:54	34.6	44.4	2.1	17.5	$ \begin{pmatrix} \bar{x} & co = \\ \bar{x} & co_2 = \\ \bar{x} & o_2 = \\ \bar{x} & o_2 = \\ \bar{x} & o_3 = \\ \bar{x} & o_4 = \\ \end{pmatrix} $
10:55	34.4	46.4	2.1	17.6	(LO MIN) { X NOX=
					-
- L	c, 2	1.0	ė. u	0.0	39tc
	44.4 7	5,4	4.0	19.0	H-SPAN
	23.1	D. 8	2.2	11.0	W-594N
	10.2				UA92-1
	43.7	75.5	2,2	19.0	BIAS

			P	Project No. Page of 8				
				Client ON1ZUKA APB				
PACIFIC ENVIRONMENTAL SERVICES, INC.				SUNNYVALE, LA				
Prepared By	Date 12-3	Checked By	Date ~	Sheet Title				

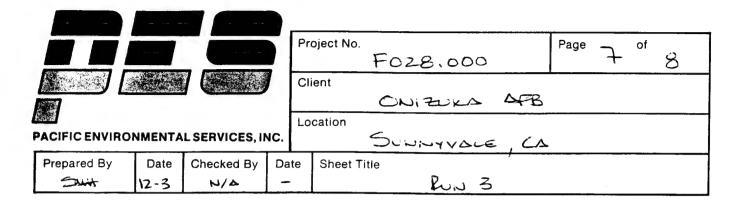
TIME	NCX	ev	COL	<u> </u>	NOTES
11:51	35,6	65.9	2,2	17.5	START TEST PLY SE PORT
11:53	35.6	ليلا، ٥	2.2	17.5	
11:31	35.6	45.8	2.1	17.5	
11:55	35.6	45.7	2.1	17.5	
11:56	35.6	65.8	2.1	17.5	
11:57	35.8	65.9	2.1	17.5	
11:58	35.8	45.8	2.1	17.5	
11:59	35 ,7	45.5	2.1	17.5	
12:00	35.6	45.5	2-1	17.5	
12:01	3 5.6	43.7	21	17.5	(lomin)
12:02	35.6	46.0	2.1	17,5	
12:05	35.6	46.0	2.2	17.5	
12:04	35.5	66.0	2.1	17.5	
12:05	35.6	66.0	2.1	17.5	
12:06	35,7	66.0	2.1	17.5	
12:07	36.0	45.9	2.1	17.5	
12:08	36.1	66.0	2.2	17.5	er e
12:04	34.0	46.0	2.1	17.5	
12:10	36.0	لوله.ع	2.1	17.5	<u></u>
12:11	36.1	66.0	2.1	17.5	(20 MIN) [MOVE to P3] *
12:12	35.5	64.5	2.1	17.5	and the second of the second o
12:13	35.3	66.7	2.1	17.5	
12:14	35.5	46.5	2,1	17.5	
12:15	35.5	46.5	2.1	17.5	
12:16	35.3	46.5	2.1	17.5	and the same of th
12:17	4 5.5	46.4 35.5	2.1	17.2	and the second second
12:18	35.9	46.4	2.1	17.5	
12:19	3 5,7	66.7	٤.١	17.5	
12:4	36.1	66,4	2.1	17.5	
12:21	36.2	نوله، ١	2.1	17.5	(30 MIN)



TIME	Nov	0	<u>cc</u> -	02	NOTES.
12:22	36-3	66.2	2.(17.5	
12:23	36.4	66.2	2.1	17.5	
12:24	36.4	46.1	2.1	17.5	
12:25	36.2	66.1	2.1	17.5	
12:26	36.2	46.4	2,1	17.5	
12:27	.36.3	46.3	2.1	17.5	
12:28	36.4	66.2	2.1	17.4	
12:29	36.2	اب <i>ات</i> ارا	2.1	17.5	
12:30	36.2	66.2	2.1	17.5	
12:31	36.1	46.2	2.1	17.5	(40 MIN) (SWITZLESS & RZ)
12:32	36.1	66.2	2.1	17.5	_
12:53	34.0	66.1	2.1	17.5	
12:34	36.0	44.0	2.1	17.5	
12135	36.1	45.9	2. (17.5	
12:36	36.(44.0	2.1	17.5	
12:34	36.1	65.9	2.1.	17.5	
12/38	36.1	45,8	2.1	17.5	
12:39	36.0	65.8	2.1	17.5	
12:40	36.0	45.9	2.1	17.5	
12:41	35.9	45, 8	2-1	17.5	(D) MID)
12:42	35.8	65.7	2.1	17.5	
12: 43	35.8	65.9	2.1	17.5	
12:44	35.8	45.8	2.1	17.5	
12: 45	35,7	65.8	2.1	17.5	
12:46	35.7	45.8	2.1	17.5	
12: 47	35.6	45.8	2.1	17.5	
12148	35.6	65.5	2.1	17.5	
12:49	35.6	45.4	2.1	17.5	
12:50	35.5	65.8	Z	13,5	
12:51	35.6	65.8	2.1	17.2	(60 MIH)

	30.127								
					Project No. Page of 6 8				
			State State of		Client ONIZUKA APB				
1				I	ocation				
PACIFIC ENVIRONMENTAL SERVICES, INC.				IC.	SUNDAVALE LA				
	Prepared By	Date	Checked By	Date	Sheet Title				
	Sum	12-3	N/A	_	Run 3				

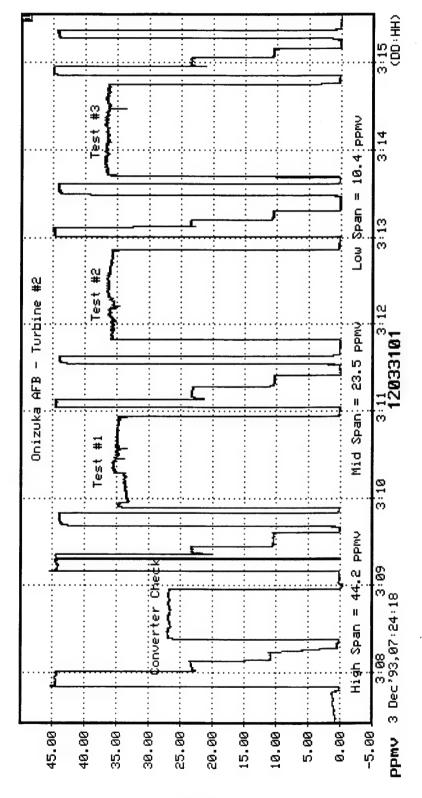
TIME	Nox	0	co-	٥٢	Notes
_ ~	0.1	0,7	-0.1	0.0	Z-20 +0.01
	44,6	75.2	3,8	19.0	H-SPAN 3.9%
مالله مورد	23.3	30.5	2.1	llio	M- 38AN
	10.3				L-SPAN
	44.0	75.0		18.9	Bias
		grand Northern Eviptum a	manner i serviciano e foreconare omerfica	2000	
			and the second s		
13:46	36.4	65.5	2,2	17,4	START AT R 2 SE
13:47	36.5	45.5	2.2	17.5	
13:48	360,60	45.5	2.2	17.5	
13:49	36.6	65.6	2.2	17.5	
13150	36.7	65.6	2.2	17.5	
13.51	36,7	65, 7	22	4 17.5	
13:52	36.6	45.6	2.2	17.5	
13:53	36.6	45. +	2,2	12.4	
13:54	36.6	45.6	2.2	12,4	
13:55	36.7	65.6	2.2	17.4	(10 MIL)
13:56	36.8	65.8	2,2	17.4	
13:57	34.6	65.5	2,2	17.4	
13:58	36 . 6	45.6	2.2	17.4	
13.59	36.6	72'Z	2.2	17.5	
14:00	36.6	65.4	2,2	17.5	
14:01	B6.5	65.3	2.2	17.5	The second secon
14:02	36.5	45.5	2.2	17.5	The second secon
14:03	36.6	45,6	2.2	17.5	
14:04	36.6	65.6	2.2	17.5	Comment of the commen
14:05	36.5	45.5	2, 2	17,5	(20 MIN) MONDTO P3 SE

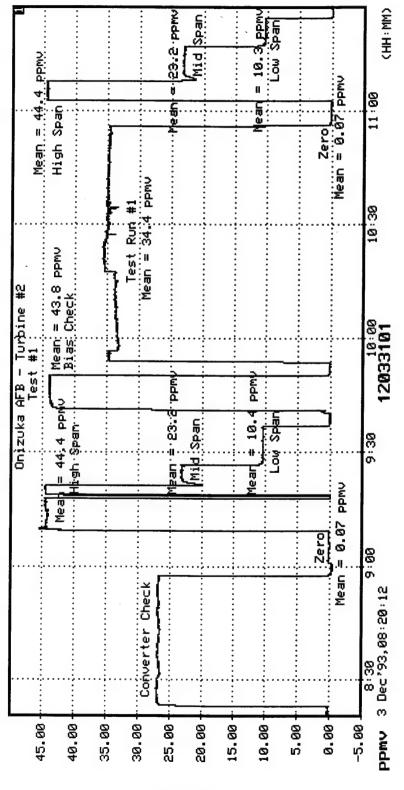


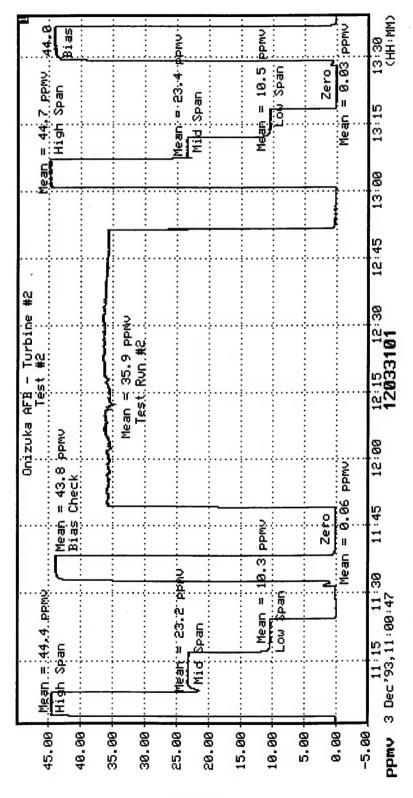
TIME	NOF	<u>co</u>	COL	0,_	HOTES:
14:06	34.3	65.5	22	17,5	
14:07	36.2	65.1	2.2	17.5	
14108	34.3	45.3	2,2	17.5	
14109	36.3	45,5	2,2	17,5	
Millo	36.3	45.6	2,2	17.5	
19:11	34.3	65.4	2.2	17.5	
14:12	36.3	65.5	2.2	14.5	
14:13	36.4	45.5	2.2	17.5	
14:14	36.5	45.4	2.2	17.5	
14:15	36.4	45.5	2.2	17.2	(30 MID)
14:16	36.3	454	2,2	17.5	
14117	36.5	45.5	2.2	17.5	
14:18	36.7	45.6	2.2	17.5	
14119	36-7	45.7	2.2	17.5	
14:20	36.5	65.8	2.2	17.5	
14:21	36.5	65.6	2.2	17,5	
14:22	36.7	65.5	2.2	17.5	
14.23	36.6	45.5	2,2	17,5	
14:24	36.6	65.5	2.2	17.5	
14:25	36.4	45.5	2.2	17.5	(40 MIN) MINED to PLY SE
14:26	36.4	65.5	2.2	4.5	
14:24	34.3	65.5	2,2	17.5	
14:28	36,2	65.5	2.2	17.5	•
14:29	36.2	45.5	2,2	17.5	
14:30	36.0	43.7	2.1	17.6	PROBE HOLED FOR PAIO
14:31	34.2	65.6	2.2	17.5	
14:32	36.0	656	2.2	17.5	
14:33	36.3	65.8	2.2	17.5	
14:37	30.4	65.6	2.2	17,5	
14:35	36.5	<i>45</i> ,8	2,2	17,5	(50 MIN)

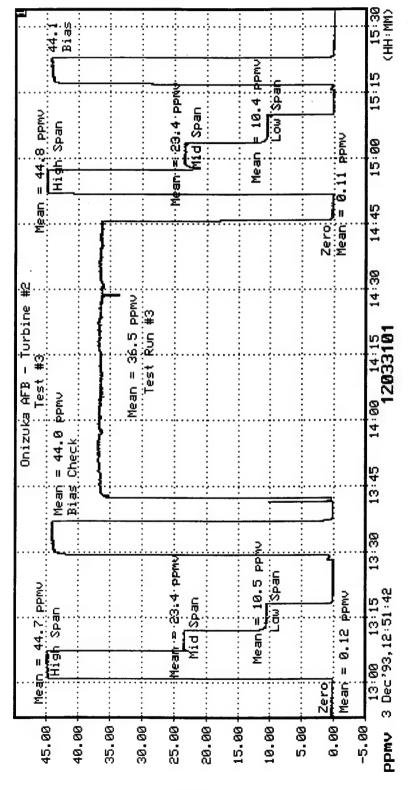
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		and the second		Project No.	028,000		Page 8	of	8
			(Client	ONIZUKA	<u> </u>	LPB		
				_ocation					
PACIFIC ENVIRO	NMENTA	L SERVICES, IN	IC.		SUNNYYA	<u>ue_</u>	, LA		
Prepared By	Date	Checked By	Date	Sheet Title			_		
Stait	12-3	10/A	_	.	V	2un	3		

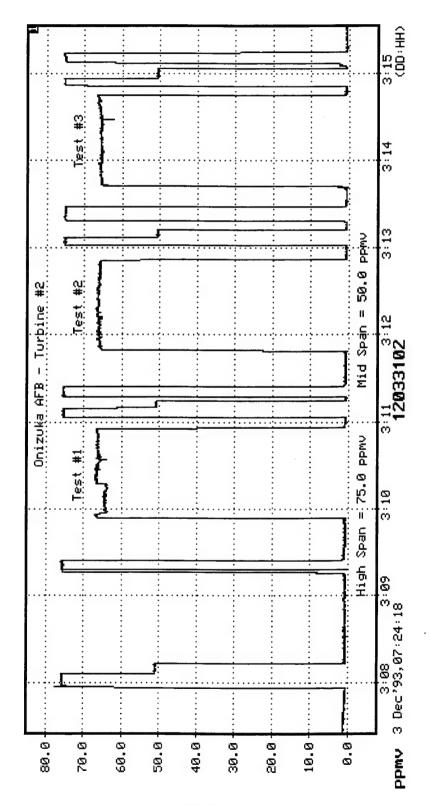
TIME	Non	Co	CO2	02	NOTES:
14:36	36.5	65.9	2.2	11,5	
(4:37	36.5	66.0	2,2	17.5	
14:38	36.4	65.8	2,2	17.5	
14139	364	66.1	2.2	12.5	
14:40	36.4	66.2	2.2	17,4	
14:41	36.2	46.3	2.2	17.5	
14:42	36,2	66.2	2.2	17.5	
14:43	30.3	66.0	2.2	17.5	
14:44	36.3	66.2	2.2	17.5	
14:45	36.2	46.4	2.2	17.5	(60 MIN)
				to the second se	
				1 1 1	
	C. Ø	0.7	0.0	6.0	25%
	44.8	75.1	3.9	19.1	U-SPAN
	23.3	50.5	2.2	11.0	M-SPAN
	10.4				L-SPAN
·	43-7	75.0	2.2	19,0	Bias
15:-	44.1				END OF TEST.
•					

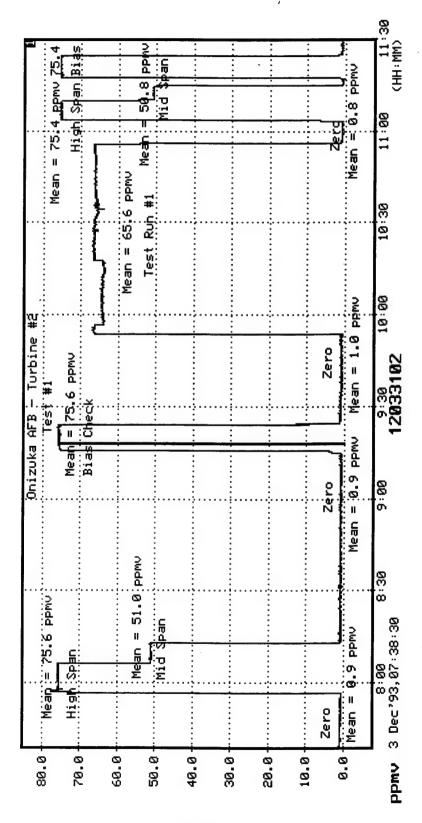


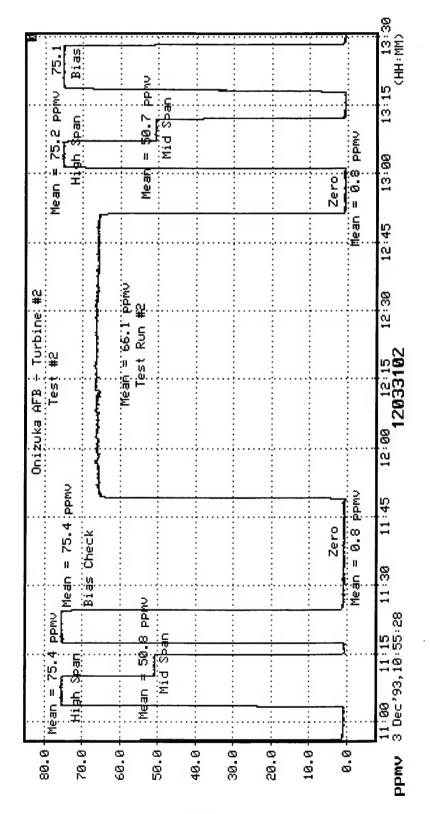


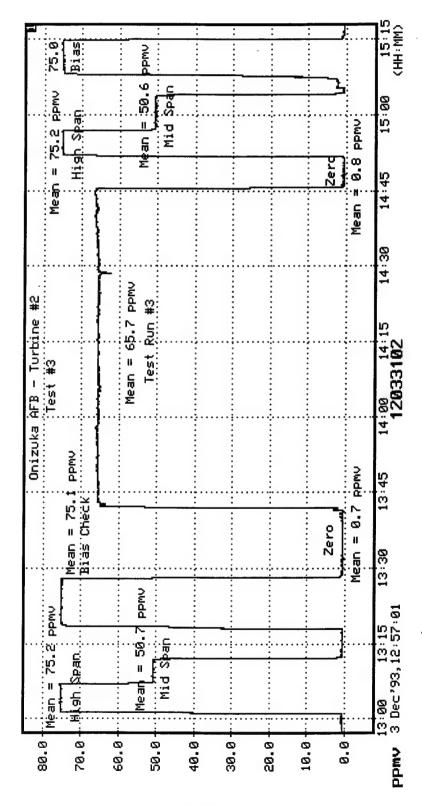


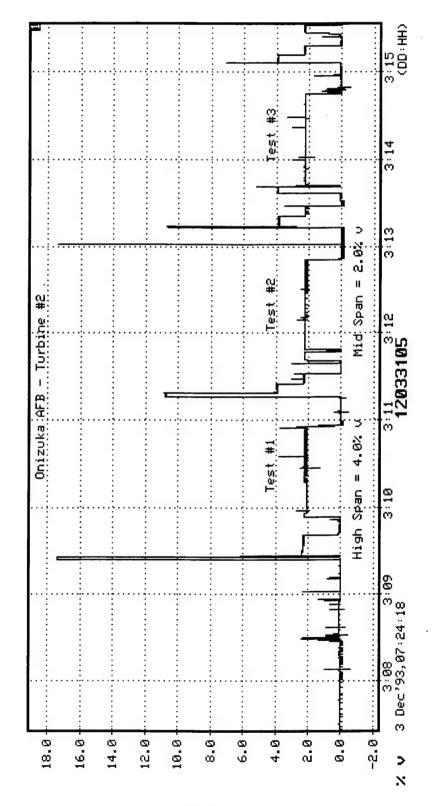


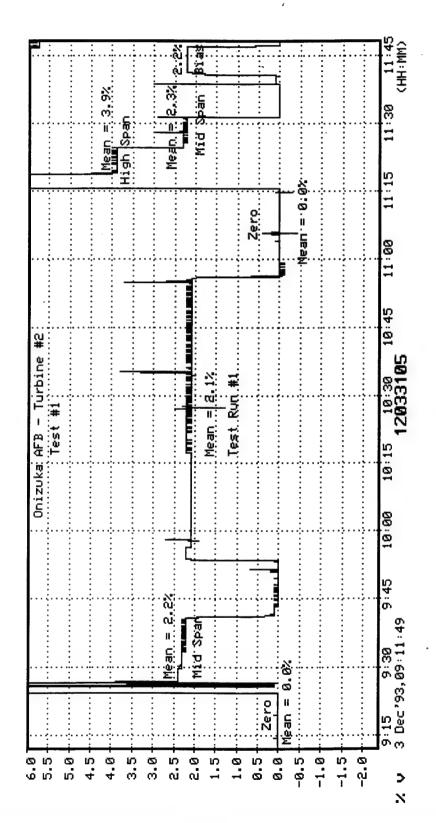


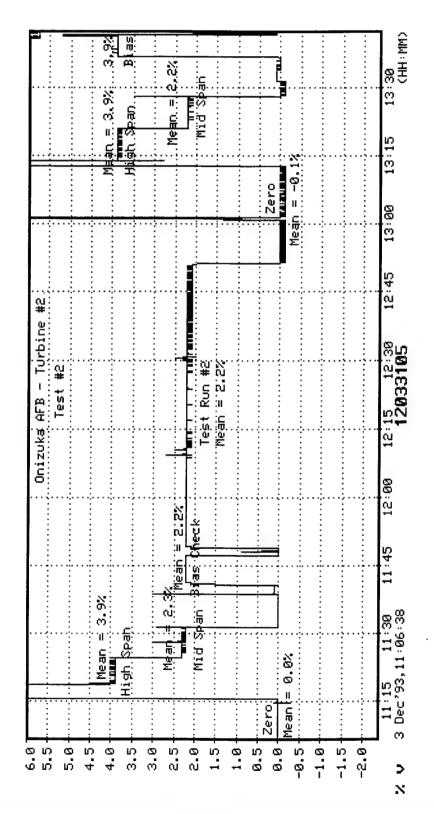


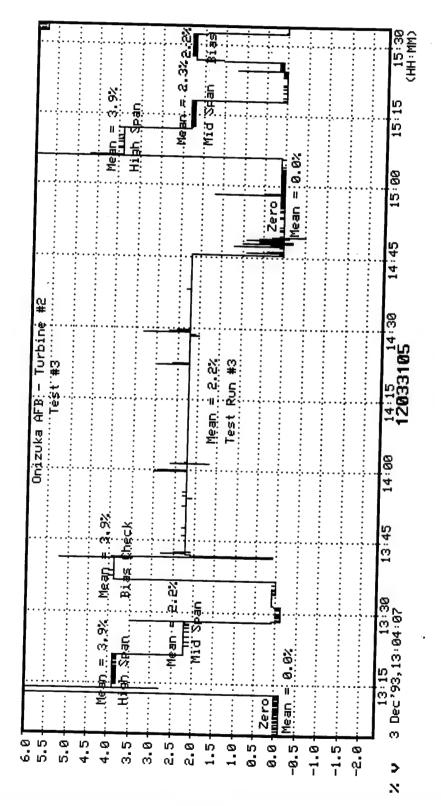


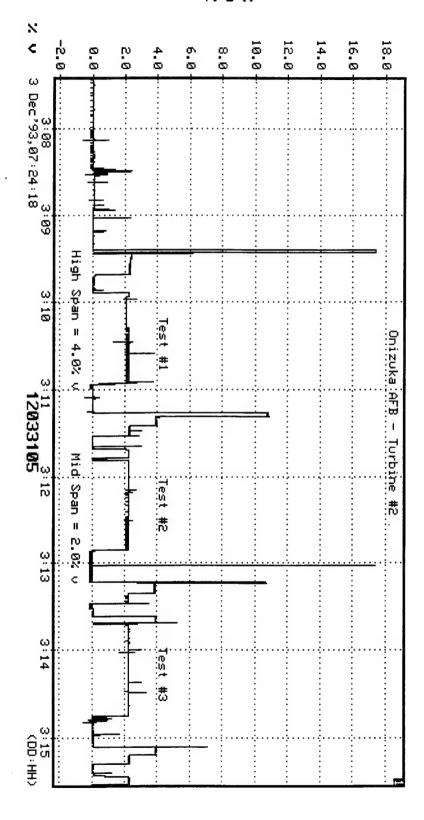


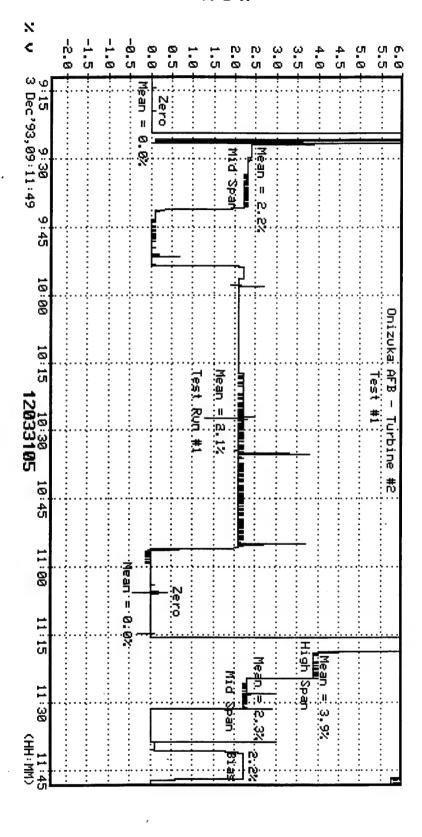


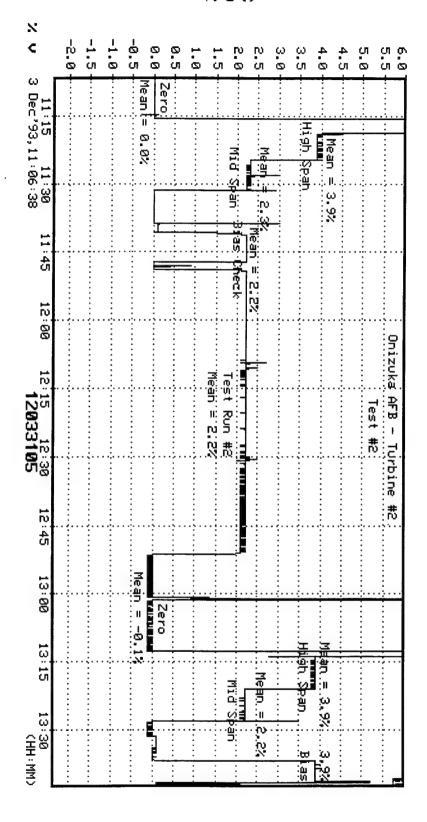


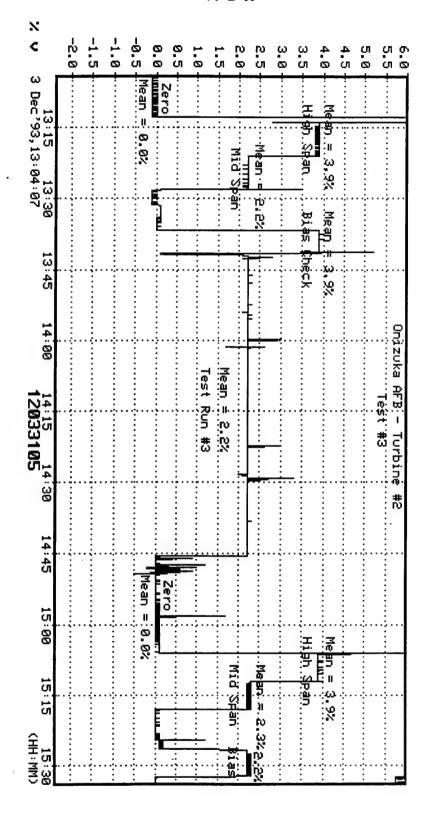


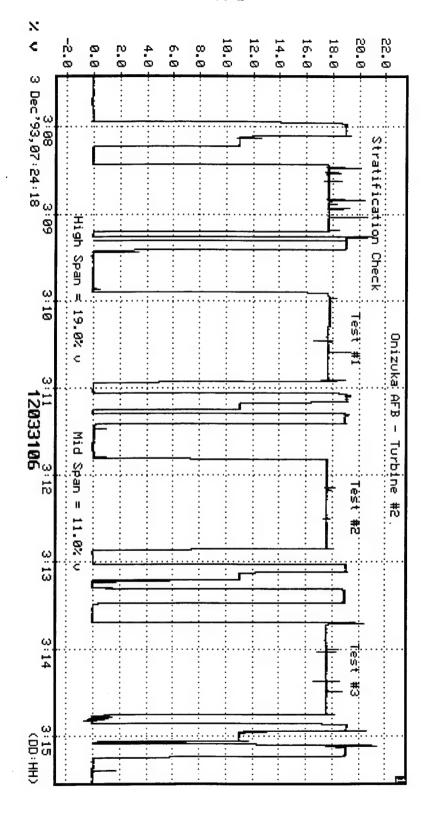


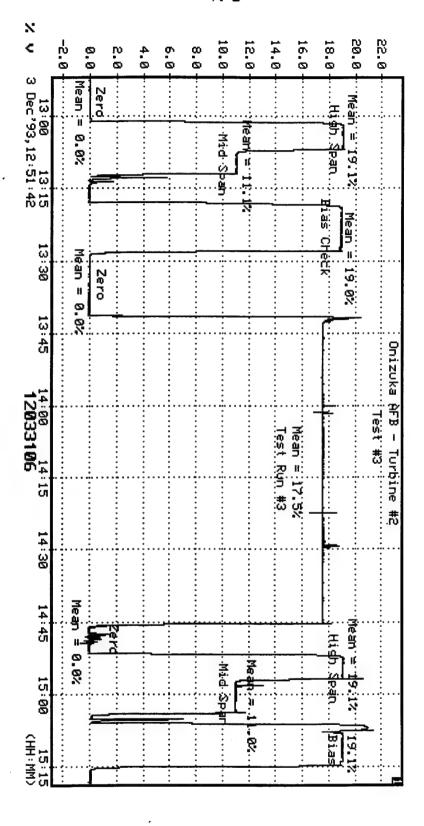














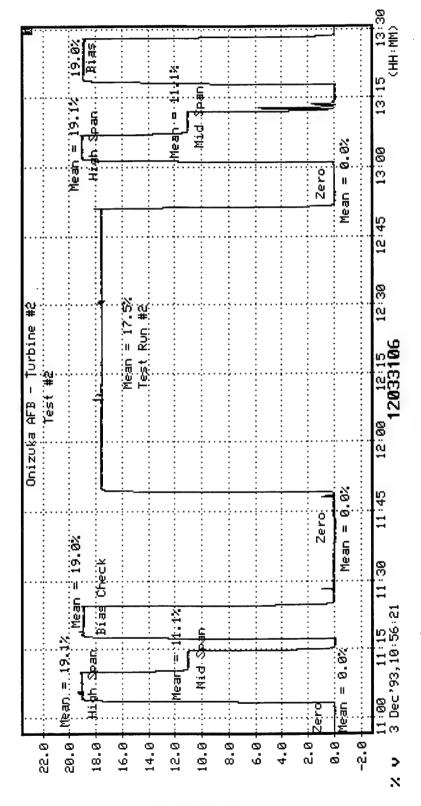
Project No.	F028		Page	ſ	of 4
Client	ONIZUKA	AFB			
Location	TIPRINE	#2 F	XHALF	T	

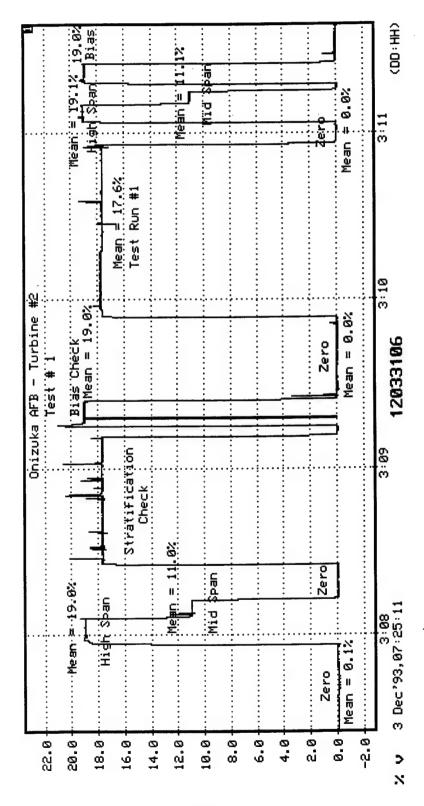
PACIFIC ENVIRONMENTAL SERVICES, INC.

Checked By Date

Sheet Title CONTINUOUS MONITORING RAJOC

1 00	/					
•		PRE -		POST-T		
NOx	SPAN VALUE PPMV	response PPMV	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR %F.S.	DRIFT %F.S.
TEST *1						
ZERO	0,00	0.07	0,14	0.07	0,14	0,00
LOW SPAN	10.4	10.4	0.00	10.3	-0.20	-0.20
MID SPAN	23,5	23.2	-0,60	23,2	-0.60	0.00
HIGH SPAN	44.2	44.4	0.40	44.4	0,40	0,00
BIAS	44.2	43.8	-0,80			
	10.100					and the second s
TEST #2						
ZERO	0.00	0.06	0.12	0.03	0.06	-0.06
LOW SPAN	10.4	10.3	-0.20	10.5	0.20	0,40
MID SPAN	23.5	23.2	-0.60	23.4	-0.20	0.40
HIGH SPAN	44.2	44.4	0.40	44.7	1.00	0.60
BIAS	44.2	43.8	-0.80			
		The address of the second seco	1			
TEST #3	Also a complex of districts of the complex of the c	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		i		
ZERO	0.00	0.12	0,24	0.11	0, 22	-0.02
LOW SPAN	10.4	10.5	0.20	10.4	0.00	-0.20
MID SPAN	23.5	23.4	- 0.20	23.4	-0.20	0.00
HIGH SPAN	44.2	44.7	1.00	44.8	1,20	0.20
BIAS	44.2	44.0	-0,40	44.1	-0.20	0,20
		•	,		gar and the second	





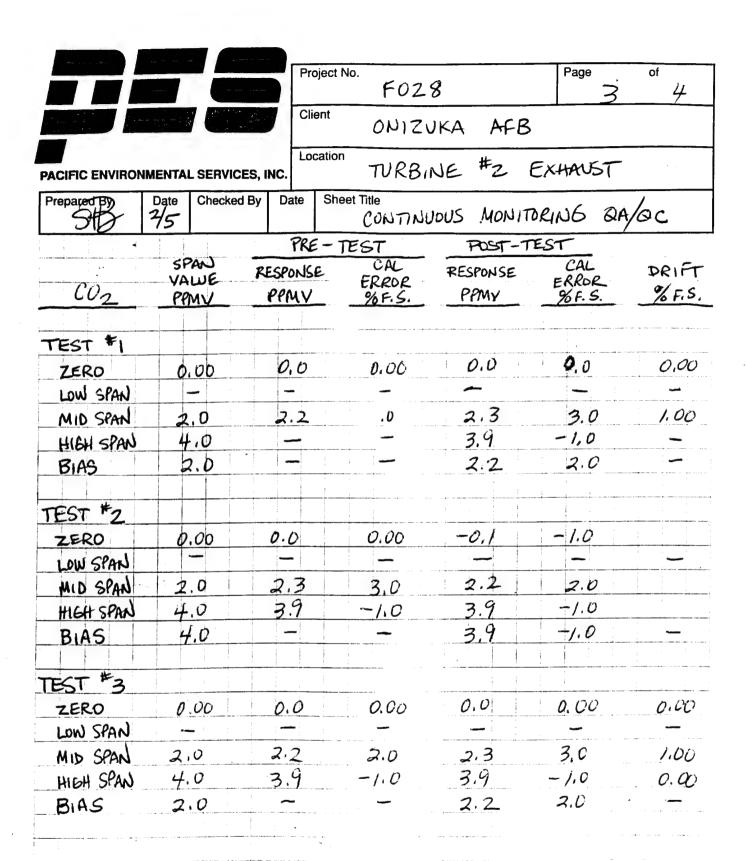


Project No	F028		Page	2	of 4	
Client	ONIZUKA	AFB				
Location	TIPRINE	#2 FX	HALK	T		

PACIFIC ENVIRONMENTAL SERVICES, INC.

Sheet Title CONTINUOUS MONITORING BAJOC

Prepared By	Date Che	cked By Date Sh	neet Title CONTNU	JOUS MONIT	dring a	a/oc
		PRE-	TEST	POST-T	EST	
CO	SPAN VALUE PPMV	PECPANCE	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR % F. S.	DRIFT %F.S.
TEST *1						
ZERO	0.00	0.9	0.90	0.8	0.80	-0.10
LOW SPAN				allowed by the control of the contro	And property consistency and a second of the	
MID SPAN	50.0	57.0	1.00	50.8	0.80	-0.20
HIGH SPAN	75.0	75.6	0.60	75,4	0.40	-0.20
BIAS	75.0	75.6	0.60	The state of the s		
					manufacture of the second of t	erragena monomonionia. Na il 1000 algano il 1000 se conditione in la conditione in the conditione in t
TEST #2			ngan kan sansan nga penggan ng 1900 - 111 - 111	· · · · · · · · · · · · · · · · · · ·		
ZERO	0.00	0.8	0.80	0.8	0.80	0.00
LOW SPAN			-			
MID SPAN	50.0	50.8	0.80	50.7	0.70	-0.10
HIGH SPAN	75.0	75.4	0.40	75.2	0.20	-0.20
BIAS	75,0	75.4	0.40			
THE CONTRACTOR			manus Milando anno marco a con a con a con			
TEST #3	e comment e i					
ZERO	0.00	0,7	0.70	0.8	0.80	0.10
LOW SPAN	<u>~</u>					
MID SPAN	50.0	50.7	0.70	50,6	0.60	- 0.10
HIGH SPAN	75.0	75.2	0.20	75.2	0.20	0.00
BIAS	75.0	75.1	0.10	75.0	0.00	





Project No.	F028	Page 4	of #
Client	ONIZUKA AFE	3	
Location	-100 10 th-		

PACIFIC ENVIRONMENTAL SERVICES, INC.

TURBINE #2 EXHAUST Sheet Title Prepared By Date Checked By Date TITLE CONTINUOUS MONITORING RA/OC 45

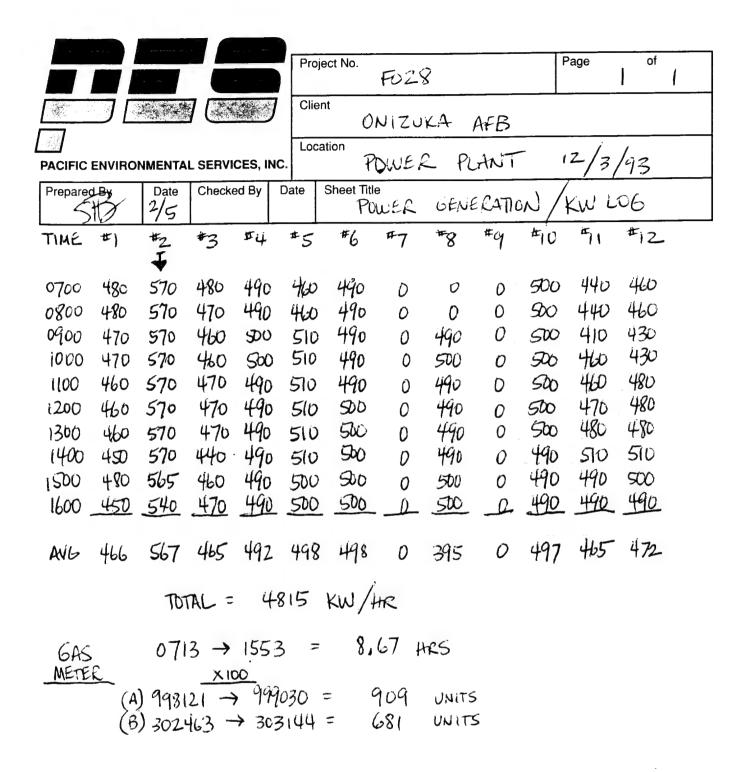
	•	PRE-	TEST	POST-T	EST	
02	SPAN VAWE PPMV	response PPMV	CAL ERROR %F.S.	RESPONSE PPMY	CAL ERROR %F.S.	DRIFT %F.S.
	· · · · · · · · · · · · · · · · · · ·					
TEST *1					1.	<u> </u>
ZERO	0.00	0.1	0.40	0,0	0.00	-0.40
LOW SPAN	-					_
MID SPAN	11.0	11.0	0.00	11.7	0,40	0.40
HIGH SPAN	19.0	19.0	0.00	19.1	0.40	0,40
BIAS	19.0	19.0	0.00	-	-	
TEST *2						
ZERO	0.00	0.0	0.00	0.0	0.00	0,00
LOW SPAN						
MID SPAN	11.0	11.1	0.40	11.1	0.40	0.00
HIGH SPAN	19.0	19.1	0.40	19.1	0.40	0,00
BIAS	19.0	19.0	0,00			
				-		
TEST #3	! -					
ZERO	0.00	0.0	0.00	0.0	0.00	0.00
LOW SPAN	_	_				
MID SPAN	11.0	11.1	0.40	11.0	0.00	-0.40
HIGH SPAN	19.0	19.1	0.40	19.1	0.40	0.00
BIAS	19.0	19.0	0.00	19.1	0.40	
		The second secon		and the second second second		The second secon

750 SPTS/DE FORM 34, JAN92 REPLACES 10048988QUDE FORM 34, OCT 89, WHICH WILL BE USED PACKAGE NUMBER TOTAL TO DATE J. C. P.S. A. 0001 2200 1400 410 700 1200 460 700 TOTAL TODAY 8 PREVIOUS TOTAL 1900 8 1700 4 60 1300 4/0 1100 460 0800 460 776 1800 \$ 33 010 000 0000 9 ENGINE OPERATING HOURS 8 0300 UST 176 0200 H/C 180 0100 11/0 18 JME. MON 11/0 ENGINE SERIAL NUMBER 337 ¥ DAILY POWER PLANTOPERATING LOG - (TURBINE - BOILER) KVAR 20 3/ca dann 124774 C. -₹-# 3 AMPRIAGATIS T-7 T-6 PCD 1084 Som SAG. NAND GENERATOR C654 630 870 69 480 cgo 40 80 21 650/8/0 69 630 810 69 630 870 45 40 820 69 ×20 068 830 63 890 82 088 63 078 0,0 25/12/12 SWINGS <u>ي</u> DAYS SQIM G 56 25 27 7,5 ç 143 6 **B** 1. 140 2 Sel 3 134 17. ē 0 ٠, Ť ٥, 63 41 63 8 TANK Son d 22 58 ر ج 7 <u>-</u> ? ij 12 3 BOILER ĭ ٦ ۲, 1700 PACKAGE NUMBER TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2300 2200 2100 2000 1800 1. 2 AT 30001 8 1500 17 30 8 38 200 <u></u> 8 8 88 0700 8 0500 2450 GENERATOR MFR ENGINE OPERATING HOURS 0300 0200 0100 N Q 3ME ENGINE SERIAL NUMBER 520 50 02 065 00 520 50 570 3 0450 1. 00 20 60 20 KVAR 60 13612 2 AMPS STARIED 20 AMPS AMPS VOLTS T-7 T-6 PCD ¥ ELECTRIC MACHINERY TURBINE AND GENERATOR . DAYS SWINGS SQIM 20646 OIL ADDED 74 700 950 740 200 B 920 00 γ 7 122 72 23 23 Z ζ. Ę ę MODEL 136 ē 'n 75 BEMAC II T-1021S-21 0 TANK MANUTAN $\widetilde{\iota_v}$ ß , Ç PACKAGE NUMBER 2300 TOTAL TO DATE TOTAL TODAY 8 2100 PREVIOUS TOTAL 8 8 1700 4/60 3/00 **1**50 8 1000 47/ 300 8 1300 470 300 1200 0500 U50 1100 420 8 0800 470 150 VOLTAGE 4160
ENGINE SERIAL NUMBER 900 0700 ENGINE OPERATING HOURS 8 8 MDN HILD 0300 HHC 75C 050 A110 520 3MIL 077 17 0010 140 300 140 300 22,000 470 460 80 480 Š D) ou 300 300 KVAR 300 290 13 ders 5 h 6 C h l - <u>A</u> 65 3 ~ À x 12 69 7240 TURBINE AND GENERATOR 93 XW 750 SQIM DAYS 20636 OIL ADDED 280 087 170.840 200 068 087 CNO 870 86 100 スプラクシ たべ 8 80 900 880 674 67 8 3 ğ 36 36 36 7 Ş 3 12 23 136 5 X Drc •6 3 63 0 53 12.315 5512712 53 7.5 15 54 17 715 12 35 ANS. 3 15 12.5 3 12.5 1 12.5 12715 2 17 5 3 16

150 SPTS/DE FORM 34, JAN 92 REPLACES 1004SPSSQ/DE FORM ACKAGE NUMBER REMARKS PUBLICATES 1000 490 20 TOTAL TO DATE TOTAL TODAY 2200 2100 8 800 PREVIOUS TOTAL 8 76 070 12 00L 1300 490 220 1200 190 220 911-617 con 1100 490 200 0900 Carl 2001 ENGINE OPER 2/1/005 0010 NGINE SERIAL NUMBER 140 m 8, 24 1 20 250 30 30 XW KVAR DAILY POWER PLANT OPERATING LOG - (TURBINE - BOILER) 125000 27 10 10 73 70 AMPS AMPS AMPS #1 URBINE AND GENERATOR 70 DAYS SÜM 106533 VOLTS T-7 T-6 PCD 2000 71136170 15 00 01 200000 200 260 1609801 5 m 080 12 22 050 H 521 65 24 240 059 12 65 44 48 34 670 830 77 67 175 680 330 75 26 SE OUT 690 82 90 360 71 2 2 DAYS SQIM 5 サンプ 57 3 57 132 53 132 S 57:19 57 57 ő 5201 0 OPERATORS ij ō 6 8 -エ 4 65 2 _ 7 τ 75 3 S 64 13 TANK N ڏ Ü 21/2 21 ù 3 2 100 PACKAGE NUMBER 1700 500 2200 2100 2000 TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2300 88 800 ğ 8 8 8 1200 . 8 1000 900 0500 9800 286 0971 0000 GENERATOR MFR
ENGINE SERIAL NUMBER 6700 0200 0100 2911 0000 NON TIME 5 00 910 NO USC 780 29,60 \$ E 85 200 120 78c 780 É KVAR 142995 TURBINE AND CI NIRATOR 24,74 # 5 ELECTRIC MACHINI RY 8 SWINGS SQIM DAYS 읃 (61) 6.00% 18. 350 V/ 901830 171 à 3 920 7 3 · ,` 2 7. 7 7 g 156 15 52 120 26 1% é MODEL 7,7 5 7 ٥٥ BEMAC II TEMP S. 7 2300 200 TOTAL TO DATE TOTAL TODAY 2100 PREVIOUS TOTAL 8 1700 UZU 1700 8 1800 1 60 (30) 8 8 200 1100 HP 1100
VOLTAGE 4160
ENGINE SERIAL NUMBER 8 0700 800 8 9 8 000 50C 700 0200 ENGINE OPERATING HOURS 0100 NDN 3 \$0 490 500 80 490 510 700 470 064 200 500 700 8 Ş Elen stens 2 Ş 180 100 8 KVAR AMPS AMPS 190 35 550 139 438 35 2014 # 0 TURBINE AND GENERATOR 00 KW 750 64522 SQIM DAYS OIL ADDED 196 00 890 80 00 600 130 890 087 1 363 CM 670 890 51 650 670 310 11 180 Soc 11/ 15: 300 10 60 35 4 1 7 7 20 20 2 > ğ ١ 'n <u>,,</u> 65 ; ζ, Ē Š 121 ₹ . 101 • •5 3 (7 62 62 10 ٤, 6. 6 3. \ _2 11 0 12 5 'n ۲-12 5 125 3 12014 125 ò 1 2 =

750 SPTS/DE FORM 34, JANEZ, EPLACES 1004SPSSQUDE FORM 34, OCT 89, WHICH WILL BE USED TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2200 200 1800 8 100 8 8 8 1200 ž 0700 800 0400 0300 Ž 3ML KW KVAR DAILY POWER PLANT OPERATING LOG - (TURBINE - BOILER) 132017 Ħ TURBINE USD GENERATOR SEC #55 The state of the state of 0612 SWINGS DAYS SGIM ę Ć 6 5 3 buce 1 უნ di TANE T BOILER REMARKS TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2300 2200 2000 900 2100 1500 (JV) 8 1700.17 0 ğ 1400 ğ 1200 Ĩ 0700 3 0900 ENGINE MFR
GENERATOR MFR
ENGINE SERIAL NUMBER 0500 800 8 0200 ENGINE OPERATING HOURS NON TIME 3 190 180 470 180 190 490 190 ¥ 170 3 180 KVAR Line & 850 Ħ 20 TURBINE AND GENERATOR 38395 20 SOLAR
ELECTRIC MACHINERY 20 SWINGS DAYS SQIM 20 m No oth 160 067 20 4 8 13 ١ 15 ę MODEL 1. 27 4 75 12 67 ٥, T-1021S-21 57 TEMP Š BOILER 6 16 PACKAGE NUMBER 2200 2000 TOTAL TO DATE TOTAL TODAY PREVIOUS TOTAL 2100 8 1800 1700 ĕ **5**00 100 300 1200 1100 8 0800 0800 0700 VOLTAGE 4160
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S AMES AMES VOLTS 7-7 1-5 PC 0 KW 750 DAYS MDS OIL ADDED ğ ē 9 w DEC •5 TEMP AIREI 83 BOILER

PACKAGE NUMBER	Flow stem that	REMARKS SWINGS	DAYS DAYS	151312	A ING HEURS THE ADDED		2200		2000	1900	(79	77147 * CO & 70 1/30 4 1/30 5	アンファング	120 120 120 120 120 120 120 120 120 120	00 870 77 57	200 200 700 80 72 57 139 P	80 120c 80 72 57 138	200 38 4 58 13	100 100 100 100 100 100 100 100 100 100	50 180 11 95 HC 068 DOC 300 SC	(5' h5 & 76 (55 h6 06) ax	5 / / 5 / / 5 5 / / 5	5/0 17c 7m 87c/7c 57 135 8 53 12.5	200 1/2 3 53 020 76 57 125 3 53 020 1	1/C De 8/0 72 37 125 \$ 54 135	30 1/0 170 174 19ch 700 860 70 57 185 8 154 127	15 170 mm 20 360 /24 57 12 - 32 56 125	CD LOP LOT LO INLET	TURBI	00000	DAILY POWER PLANT OPERATING LOG - (TURBINE - BOILER) ENGINE SERIAL NUMBER
PACKAGE NUMBER # // //	REMARKS Und boung down \$7.0	TOTAL TO DATE SWINGS	TOTAL TODAY DAYS	2157 "	ENGINE OPERATING HOURS OIL ADDED	2300		2002		1800	100 1/ W 1/71 (O1	1 400 le	17	100 510 110 110 110 110 110 110 110 110	100 4 061 45 06 016 000	420 180	1 E1 67 h OAT 65 06 018 080	80 20 54 13 4 65 13 15 15 15 15 15 15 15 15 15 15 15 15 15	120 64 63 06	#0 200 /20 /20 /20 /20 /20 /20 /20 /20 /2		12 C 177 17 52 (1/2 1/2 cmg ch)	94 1 1 5 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	100 Mill 19 1/2 1/2 1/2 1/2 1/2 1/2 1/2 1/2 1/2 1/2	27 h 581 AS 14 329	5-11 CT 11-561 115 11. 11. 1009 89 19-59 60:	650 1/ 1/2 0 034 V (1) 7 4	TIME KW KVAR AMPS AMPS AMPS AMPS AMPS POLITY TO LOP LOT LO MILE PS	1/0/24	JUMBER	SOLAR MODEL MODEL
PACKAGE NUMBER # /2	REMARKS Clan dian 1 4660 73/	TOTAL TO DATE SWINGS	DAYS	MEVIOUS TOTAL 1)62 % MIDS 8 CALL \$ 0050	GINE OPERATING HOURS	2000	2100	:000	580		19 0 (5) (1,00 to 30 me 6) 30 me 64 mil	50, 20 17 ,50	200 200 2012	20 70 1 57 ng 1 11 3	130 July 200 24 1 20 1 20 1 20 1 20 1 20 1 20 1	1200 480 200	1100 430 200	130 180 01 61 02 680 850 71	51 15 1 10 11 25 26 000 060 11 11 11 11 11 11 11 12 12 12 12 12 12	9 0	0000	050 75 4 98 069 051 05H	400 830 77 ST 170 B 50 175	C 4. 22 C 22 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	0200 1.50 1.50 1.50 0.50 0.50 0.50 1.50 1.	0.000 1.00 1.00 1.00 1.00 1.00 1.00 1.0	MON HUGA 1/2	TURBINE AND GENERATOR	I.	L	



4815 KWH \times 8.67 HRS = 41746 KWH 41746 KWH / 909 UNITS = 0.0218 WITS/KWH 41746 KWH / 681 UNITS = 0.0163 UNITS/KWH

PRELIMINARY VELOCITY TRAVERSE

Plant:	ONIZUKA AFB
Date: _	1-12-94
Location	
Stack I	
	ric Pressure, in. Hg: 30,22
Stack G	auge Pressure, in. H20:6./
Operato	
	ube I.D. Number: ST4
Tempera	ture Readout I.D.: FISHER/TC-12
Ditot To	the Leak Check: OK



Schematic of Traverse Point Layout

MABY:	CA 59	
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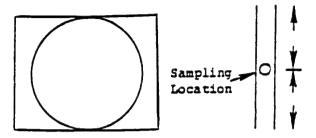
MAGY	: CA 59_		
	Velocity	Stack	Cyclonic
Point	Head (Δp_g)	Temp.	Flow Check
Number	in. H ₂ O	(Tq), °F	o from Null
L 1	1.40	450	1.6
2	1.25	451	6.0
3	1.60	438	0,8)
4	1.25	434	22.3
		443	
Average			

Traverse	Velocity	Stack	Cyclonic
Point	Head (Apg)	Temp.	Flow Check
Number	in. H ₂ O		° from Null
RI	1.40	444	
2	1.45	450	
3	1.55	449 444	
4	1.25	444	
		447	
	·		
	V		
Average	1,18	445	

- PACIFIC ENVIRONMENTAL SERVICES, INC. -

PRELIMINARY VELOCITY TRAVERSE

Plant:	ONIZUKA AFB
Date:	1-12-94
Location:	TURBINE #2 STACK
Stack I.D	24"
Barometri	Pressure, in. Hg: 30,22
Stack Gau	ge Pressure, in. H2O:
Operators	BROWN
Pitot Tub	I.D. Number: ST4
Temperatu	e Readout I.D.: FISHER/TC-12
Pitot Tub	Leak Check:



Schematic of Traverse Point Layout

MAG	1 = CA 59		
Traverse	1	Stack	Cyclonic
Point	Head (Δp_g)	Temp.	Flow Check
Mumber	in. H ₂ O	(T _q), °F	° from Null
REF.	POINT	L-3	
TEST	#1		
10:55	1.65	440	
11:05	1.65	441	
11:15	1.65	442	
11:25	1.65	441	
11:35	1.65	443	
TEST	*2		
12:19	1.65	444	
12:29	1.65	442	
12:39	1.65	444	
12:49	1.65	445	
12:59	1.65	444	
·			
Average			

Traverse Point Number	Velocity Head (Δp_g) in. H ₂ O	Stack Temp. (T _q), °F	Cyclonic Flow Check * from Null
Mumber	III - 1170	1.4//	
TEST	[#] 3		
13:38	1.65	444	
13:48	1.65	445	
13:58	1.65	445	
14:08	1.65	444	
14:18	1.65	445	
			·
Average			

PACIFIC ENVIRONMENTAL SERVICES, INC. -



HYDROCARBON SAMPLING FIELD DATA

Project No. F028

CLIENT: ONIZUKA	AFB		•	Date:	1-12-94
Sampling Location: TANK # # : FLOW METER; Time		TURBINE 108 6F99 Sample A	. 1	CHAUST 117 MP 41 ple B	10 6 6F99 Sample C
<u> </u>		" Hg "140 30.5 0.80	12: <u>18 3</u>	Hg	" Hg "H ₂ 0
<u>10</u> 20		2 5 .0 0.90		<u>5.2 0.</u> 98 0.2 <u>1.</u> 0	13: <u>48 24.8 0.90</u> 13:58 19.2 0.90
30		12.8 0.84			14:08 14.2 0.90
40	11:36	7.5 0.88	12:58 1	0.2 1.0	14:18 8.7 0.90
POST LEAK-CHECK					
0	12:22			10.4 0.00	
10	12.32	7.6 0,00	13:50	10.4 0.00	14: <u>48 8,5 0,</u> 00
		oK_	(OK	OK
		TEST #1	TES	ST #2	TEST #3
Pre Leak-checkOY			Pos	st Leak-	check

PACIFIC ENVIRONMENTAL SERVICES, INC. -



Plant: <u>ONIZUKA AFB</u>

Date: <u>1-12-94</u>

Source/Sample Number: TURBINE #2 EXHAUST RUNS 1,2.3

1. $Vm(std) = (17.64)(Vm)(Y) \left[\frac{P_{bar} + (\triangle H/13.6)}{Tm} \right]$ Vm(std) = (17.64)(Vm(std) = NA dscf.

2. Volume water vapor collected (standard conditions).

V(lo) = _____ condensate from impingers and selica gel.

Vw(std) = (0.04707) V(10) = (0.04707)(

Vw(std) = NA scf.

3. Percent moisture, by volume.

 $Bw_{S} = \frac{Vw(std)}{Vw(std) + Vm(std)} = \frac{4.6\%}{() + ()} = \frac{4.6\%}{()}$ Bws = 0.046 . * FROM PMIO TESTING

4. Molecular weight, stack gas.

Dry molecular weight.

 $Md = 0.440(% CO_2) + 0.320(% O_2) + 0.280(% N_2 + % CO)$

Md = 0.440 (2.5) + 0.320(17.5) + 0.280(80.0)

Md = 29.00 1b/1b-mole.

 $Ms = Md + Bw_s (18 - Md) = (29.10) + (0.046)(18 - 29.10)$

Ms = 28.59 lb/lb-mole.



Plant: ONIZUKA AFB

Date: 1-12-94

Source/Sample Number: TURBINE #2 EXHAUST

5. Stack gas velocity average.

$$Vs(avg) = (85.49)(Cp)(V\Delta P) \left[avg \sqrt{\frac{(Ts)}{(Ps)(Ms)}} \right]$$

$$Vs(avg) = (85.49)(1.00)(1.18) \sqrt{\frac{(460+445)}{(30.22)(28.59)}}$$

$$Vs(avg) = 103.2 \text{ ft/sec.}$$

6. Stack volumetric flow rate, actual conditions (stack temperature and pressure).

Qs =
$$(60)(Vs)(A) = (60)(i03,2)(3./4)$$

Qs = $(60)(Vs)(A) = (60)(i03,2)(3./4)$

7. Stack volumetric flow rate, standard conditions (68 degrees F, 29.92 Hg).

Q(std) =
$$(17.64)(Qs)(1-Bw_s) \frac{(Ps)}{(Ts)}$$

Q(std) = $(17.64)((9,450)(1-0.046)) \frac{(30.22)}{(905)}$
Q(std) = $(0,930)$ dscfm.

8. Isokinetic variation.

$$z_{I} = (K) \left[\frac{(T_{S})(V_{m}(std))}{(P_{S})(V_{S})(A_{n})(\theta)(1 - Bw_{S})} \right]$$

$$z_{I} = (0.0945) \left[\frac{()}{()}()()()(1 - Bw_{S})(1 - Bw_$$

ENGINE MFR SOLAR												MODEL 7-1021S			21	-21		
GENERATOR MFR ELECTRIC A							ACHINERY				MODE	EL	BEN	MAC N				
ENG	HE SE	RIAL N	UMBER	•			2	Ø	64	16								
				TI	URB	INE	AND		RATO						BOIL	ER.		
TIME	KW	KVAR	AMPS	AMPS	AM	-5 V	OLTS	Ť-7	7-5	PCD	LOP	LOT	LO	TEMP	PSI	₽		
MON					mi													
0100			25		1111	1113												
						TI I								-				
0200																		
0300			-30															
0400			1. ž															
0500			6.2															
0600			-5															
0700																		
0600			1															
0800	€60	-	75	74	7.	>		GN	870	13	57	1360	5	52	2.5	15		
1000	550				H			BU	880	15	57	1260	હ	56	12.5	7		
	550							3		745	57	130	3	-	12.5	_		
	-		W. 100						-	-				100	10.3			
1200	c .							7	0.00	4	57	136	5	64	12.5			
1300	550	_			Ш			3	900		-	-			15.2	15		
1400	221				Щ			700	900	72	57	136	5	66	-			
1500	540		200		Ш	Ш			905		58	13/0	5	68	125	15		
1000	540							710	905	72	58	12	5	68	12.5	15		
1700																		
1800									20	a	re	A.	b	177	15			
1900			1.0									-	1					
2000		-	1						1									
2100	-										T							
	┼─	+-							1		1		1	1				
2200	-	+-							+	-	+	-	1	+	1	T		
2300	1	One	RATIN	C HU			C	L ADI	DED.	-		1	<u></u>	1	L.,			
	vious						MIC											
PRE	V 1.005	IUIA	1/5	520				-		-								
TOT	AL YO	DAY					DAY											
тот	AL TO	DATE	15	ુર દ	0	7	SWM	ıG5										
REN	ARKS					_		-										

1/12/94

PACKAGE NUMBER

d



APPENDIX C LABORATORY REPORTS AND ANALYTICAL METHODS

- PACIFIC ENVIRONMENTAL SERVICES, INC. -

REPORT

TRUESDAIL LABORATORIES, INC.

CHEMISTS - MICROBIOLOGISTS - ENGINEERS

RESEARCH - DEVELOPMENT - TESTIN

PACIFIC ENVIRONMENTAL SERVICES, INC.

CLIENT Baldwin Park, CA 9

Baldwin Park, CA 91706 Attn: S. Hugh Brown

SAMPLE

3 tanks from project PES/ONIZUKA AFB

R R

14201 FRANKLIN AVENUE TUSTIN, CALIF. 92680 AREA CODE 714 • 730-6239

AREA CODE 213 • 225-1564 FAX 714 • 730-6462

Jan. 18, 1994

DATE

RECEIVED Jan. 14, 1994

LABORATORY NO. 53691

INVESTIGATION

Total hydrocarbon analysis by SCAQMD method 25.2

RESULTS

The submitted samples were analyzed for $\mathrm{CH_4}$, CO , $\mathrm{CO_2}$ and nonmethane hydrocarbons (as $\mathrm{C_1}$) by SCAQMD Method 25.2. Oxygen concentrations were determined by Orsat analysis.

The results obtained are as follows:

This report applies only to the sample or samples investigated and is not necessarily indicative of the quality or condition of apparently identical or similar products. As a mutual protection to clients, the public and these Laboratories, this report is submitted and accepted for the exclusive use of the client to whom it is addressed and upon the condition that it is not to be used, in whole or in part, in any advertising or publicity matter without prior written authorization from these Laboratories.

PACIFIC ENVIRONMENTAL SERVICES, INC. LN 53691

SAMPLE	<u>ID</u>	NMVHC ppmvC ₁	CH ₄	CO ppmv	CO ₂	0 ₂ %V
Test-1 Test-2	108 117	ND 11	7 6	60 58	18610 18590	17.9 18.2
Test-3	106	ND	6	57	18700	18.1
Detection	limit	4	2	2	2	0.2

ND = Not detect

TRUESDAIL LABORATORIES, INC.

Prepared by:

Xuan Huong Dang Analytical Chemist

Air Pollution Testing

Reviewed by:

Charles M. Figueroa Project Manager Air Pollution Testing



TCA-SOP Revision: Draft Date: 11/91

Prepared By: Page 1 of 10

DETERMINATION OF HYDROCARBON EMISSIONS BY TOTAL COMBUSTION ANALYSIS (TCA) METHOD

Table of Contents

- 1.0 Applicability
- 2.0 Equipment List
- 3.0 Preparation of sampling train
- 4.0 Sampling at test site 5.0 Analytical procedure
- 6.0 Gaseous sample analysis
- 7.0 Condensate trap recovery 8.0 Calculations
- 9.0 Diagrams

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Method

This method is based on SCAQMD METHOD 25.1

1.0 Summary of Method

This procedure uses a sampling train comprising a stainless steel probe and a freeze-out trap connected to an evacuated seven-liter tank via a Magnehelic pressure differential gauge. The trap is used for collecting the condensable organic matter while the non-condensable gases are being collected in the tank.

2.0 Equipment list

- 2.1 7-liter tank with vacuum gauge
- 2.2 Stainless steel condensate trap and probe
- 2.3 Magnehelic gauge
- 2.4 Metal Dewar flask with dry ice
- 2.5 Tank holder

3.0 Preparation of sampling train

- 3.1 Determine the number of tanks and traps required, and match only outlet tanks with outlet traps for sampling at an outlet location. Similarly, the principle applies to inlets.
- 3.2 Using a high-volume vacuum pump, evacuate the 7-liter tanks to a pressure of 1 Torr or less three times, filling the tanks to one atmosphere between evacuations.
- 3.3 After the third evacuation, turn off the valves and check for leaks by allowing the tanks to stand for at least 16 hours, after which period any leaks will become apparent by a change in the vacuum gauge readings.
- 3.4 For convenience in transporting as well as ease of handling at the test site, two tanks can be placed in a wooden holder.

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- 3.5 Set up each train in the following sequence: tank, Magnehelic gauge, trap, and probe. Be sure to determine the proper torque required to tighten these crucial connections, as too much torque will spoil the fittings for future conections while too little torque will result in leakage.
- 3.6 Perform a leak check on each train by introducing nitrogen into the connections at the probe tip. Magnehelic needle will deflect and should return to the same "zero" point after a short while. This signifies a good leak check.
- 3.7 The assembly is now ready for transport to the test site. It should be noted that triplicate samples are usually taken at the inlet locations and duplicates for outlets.

4.0 Sampling at test site

- 4.1 Conduct pre-test flow and temperature measurements according to method described in Truesdail Laboratories, Inc. S.O.P. Before the stack traverse, make sure that both the air pollution control unit and the production line is running at the conditions in the test protocol or on the AQMD's "Permit to Construct."
- 4.2 Immerse the traps into the metal Dewar flask filled with crushed dry ice to a depth of four inches. Maintain the dry ice level for the whole duration of the test.
- 4.3 Uncap the probe tips and wrap the tips as close as possible to each other in order to satisfy duplicate or triplicate sampling. Avoid touching the tips to prevent contamination.
- 4.4 Insert the probes into the sampling port carefully so as not to scrape the port walls on the way in. Position the tips near the center of the duct. The port should then be sealed off with duct tape.
- 4.5 Coordinate all sampling stations, i.e., begin and end the test simultaneously. Integration times will vary from 30 to 120 minutes.
- 4.6 During the sampling period, record the vacuum gauge reading at 5-minute intervals and make field notes regarding any unusual events which may affect subsequent analytical results. Examples being: (1) the plugging of flow due to a frozen line, and (2) problems associated with the production line.

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- 4.7 Adjust the flow into the tanks such that between 5 to 10 inches of vacuum remain at the end of the sampling period. Close all valves at the end of the test.
- 4.8 Remove the probes from the duct carefully and cap off the probe tips.
- 4.9 Perform a post-test leak check by opening the valves. The Magnehelic needle will deflect and will settle down to its original position if there are no leaks. Make a note otherwise.
- 4.10 Label all tanks and traps accordingly, i.e., include the name of client, sampling date and location, and test and tank number.
- 4.11 For verification and quality control purposes, conduct a post-test flow and temperature measurement.
- 4.12 Transport the assemblies to the laboratory for analysis.
- 4.13 Disconnect the sampling trains and plug the open end of the traps. Store the traps in a dry ice or household freezer until they can be processed.
- 4.14 Measure the pressure in the tanks with a manometer and add pre-purified dry nitrogen to an absolute pressure of at least 860 Torr.
- 4.15 Record these pressures as they will be needed for further analytical computations. The corrected barometric pressure and temperature are to be recorded as well.

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5.0 Analytical description

This involves the separate determinations of carbon monoxide, methane and carbon dioxide, and the combined determination of C_2 and higher molecular weight hydrocarbons. Results are reported as parts per million (ppm) and pounds per hour (lb/hr) as carbon.

The gaseous portion and condensables portion are analyzed separately. The analysis of the gaseous portion requires a gas chromatographic column to separate and elute, in order, carbon monoxide, methane and carbon dioxide in the sample. The separated components are then methanized and detected by a flame ionization detector (FID). The amount of methane measured by the FID is recorded on a chromatogram strip chart.

The analysis of the volatile hydrocarbons in the gaseous portion requires a gas chromatographic column preceded by a 7-inch loop of Tenax material placed in an ice bath (0°C) which absorbs the C₃ and higher hydrocarbons. The C₂ hydrocarbons are separated from carbon monoxide, methane and carbon dioxide in the sample (which elute together) by the chromatographic column. After C₂ is eluted, the carrier gas flow direction is reversed and the Tenax loop is heated with boiling water (100°C) to desorb the remaining hydrocarbons. As each component is eluted, it passes through a catalytic oxidizer which converts it to carbon dioxide. Each carbon dioxide peak is measured by a non-dispersive infrared (NDIR) spectrophotometer, utilizing a carbon dioxide detector, and quantified by a computing integrator (GC/NDIR).

The condensable portion of the sample is analyzed for total hydrocarbons as carbon by volatilizing the trap contents and catalytically oxidizing everything to carbon dioxide which is then collected in an evacuated vessel and quantitatively determined by the FID as mentioned in 5.2.

In order to obtain meaningful analytical data, it is necessary to procure accurate reference standards and to calibrate the instruments with these standards at frequent intervals. Known concentrations of carbon monoxide, methane, carbon dioxide and propane in dry nitrogen are purchased from a vendor and are NBS traceable (+/-2%).

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6.0 Gaseous sample analysis

- 6.1 Record the room temperature and barometric pressure.
- 6.2 After the sample tanks have settled down to room temperature, the absolute pressure is measured with a mercury manometer. The tanks are then pressurized with dry nitrogen to at least 32 inches of mercury (absolute), re-equilibrated, and measured again. The measurements are recorded and a dilution factor calculated.
- 6.3 The GC/FID instrument is calibrated with a standard gas sample.
- 6.4 The pressurized sample is used to flush the sample loop on the gas chromatograph with sample and the injection valve actuated to place the loop into the carrier circuit.
- 6.5 The sample fractions are eluted in order methane, carbon monoxide, and carbon dioxide and plotted on a strip chart chromatogram.
- 6.6 A computer instantaneously integrates and calculates each sample peak with the appropriate constants and correction factors, and reports these values at the end of the chromatogram. Replicate runs are made until \pm 5% maximum deviation is obtained.
- 6.7 The GC/NDIR instrument is standardized with a standard gas sample.
- 6.8 The pressurized sample is used to flush the sample loop on the gas chromatograph with sample.
- 6.9 The injection valve is actuated, placing the sample loop into the carrier circuit with the Tenax loop immersed in an ice bath.
- 6.10 When the C_2 hydrocarbons have been eluted or their retention time passed, the carrier gas flow through the Tenax and GC column is reversed and the Tenax loop immersed in boiling water. The C_3 and higher molecular weight hydrocarbons are eluted together and plotted with the previous peaks on a strip chart chromatogram after detection by the NDIR analyzer.

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- 6.11 A computer instantaneously integrates and calculates each sample peak with the appropriate constants and correction factors and reports these values at the end of the chromatogram. Replicate runs are made until ± 5% maximum deviation is achieved.
- 6.12 In the event that any of the hydrocarbon peaks are high enough to make the analyzer off scale, the sample is reanalyzed using a smaller loop and an appropriate standard.

7.0 Condensate trap recovery

- 7.1 Place the trap in a dry ice cooling bath and then heat the ends of the trap with a Bunsen burner to drive the hydrocarbons into the cold section of the trap away from the plugs.
- 7.2 After a minimum of five minutes in the cooling bath, the plugs are removed from the trap and the trap is then connected to the carrier gas on one end and a 1.8-liter evacuated vessel on the other end. The trap is purged at a rate similar to the sampling rate until at least 1 liter of purged gases are collected for at least six minutes.
- 7.3 Remove the purge gas vessel. Another evacuated collection vessel is the attached to the NDIR effluent and flow re-established in a push-pull fashion. The trap is connected to the oxidizer, followed by a sulfuric acid bubbler to remove moisture and a NDIR carbon dioxide analyzer to indicate how much hydrocarbon is left in the trap.
- 7.4 The cooling bath is removed and the trap is slowly heated with a Bunsen burner until the stainless steel trap reaches a dull red glow and the sample is eluted from the trap as indicated by the NDIR.
- 7.5 The collection vessel is analyzed for carbon dioxide by GC/FID as in 6.1 thru 6.6.
- 7.6 The purge gas vessel is analyzed for volatile hydrocarbons by GC/NDIR as in 6.7 thru 6.12.

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7.7 For traps where low condensables are expected, especially outlet ones, with the possibility of high moisture and carbon dioxide in the samples, the trap is placed in a water bath at room temperature and connected to a short loop of stainless steel (1/4-inch 0.D.) packed with quartz wool and placed in a dry ice bath. The purge step is then performed and the purge loop incorporated into the trap analysis as in 7.3. This alternative procedure will minimize the amount of unpurged stack carbon dioxide which may have dissolved in the trap condensate.

8.0 Calculations

- 8.1 Condensable hydrocarbons:
 - (a) Integrate the area of the standard.
 - (b) Integrate the area of the sample.
 - (c) Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smp1} = \frac{C_{std} \times A_{smp1} \times V_{ves} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}{A_{std} \times V_{tank} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}$$

where C_{smpl} = concentration of the sample in ppm, C_{std} = concentration of the standard in ppm, A_{smpl} = area of the sample, A_{std} = area of the standard, V_{tank} = volume of the sample tank in liter, V_{ves} = volume of the collection vessel in liter, P_{tank} = pressure (absolute) of the tank in inches of mercury, and T = room temperature in $^{\circ}F$.

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- 8.2 Volatile hydrocarbons and gaseous components:
 - (a) Integrate the area of the standard.
 - (b) Integrate the area of the standard components.
 - (c) Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smpl} = \frac{C_{std} \times A_{smpl} \times \frac{P_B + P_f}{P_B + P_i}}{A_{std}}$$

where P_B = barometric pressure (net) in inches of mercury,

P_i = residual pressure of sample tank, and
P_f = final pressure of sample tank after N₂
addition.

8.3 The following formula is used for the computation of the emission rate in lb/hr carbon:

$$\frac{\textit{C} \times 12 \; \textit{lb/mole} \times \textit{Q}_{\textit{sd}}}{3.79 \; \times 10^8 \; \textit{ft}^3/\textit{lb.mole}}$$

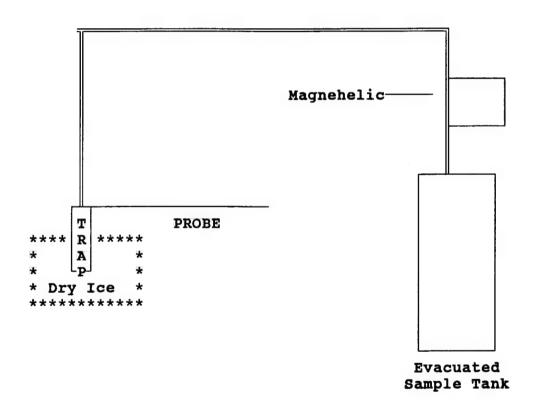
where C = total concentration of hydrocarbons present in ppm, and Q_{sd} = flow rate in standard ft³/hr (dry).

9.0 Diagrams

Diagrams of the assembly is shown on the following page. Note that the distance between the dry ice top surface and the sampling probe of the trap should be at least one inch so that water will not freeze in the narrow sampling tube.

Hydrocarbon - SOP Revision: Page 9 of 10

TCA Sampling Train Setup



Revision: Draft

Date: 11/91
Prepared by: XHD
Approved by: PDM
QA Approval: TJP
Page: 1 of 8

STANDARD OPERATING PROCEDURE FOR: TOTAL HYDROCARBON ANALYSIS BY SCAQMD 25.1

Table of Contents

- 1.0 Method
- 2.0 Principle
 3.0 Applicability
 4.0 Apparatus
 5.0 Reagents
 6.0 Procedure
- 7.0 Calculations

25.1/SOP Revision: Draft Page: 2 of 8

STANDARD OPERATING PROCEDURE FOR: TOTAL HYDROCARBON ANALYSIS BY SCAQMD 25.1

Method

This method is based on SCAQMD METHOD 25.1. It is intended for the determination of total hydrocarbons in source emissions.

1.0 Principle

Samples are collected in two fractions by using an evacuated tank to draw gases through a condensate trap chilled in dry ice. In this modification of the method, both fractions are separately analyzed for carbon monoxide, methane and carbon dioxide, in addition to a combined determination of C2 and higher molecular weight hydrocarbons. Results are reported as parts per million (PPM) in the sample, and as pounds of carbon per hour from the source.

Applicability

This method is applicable to determination of stationary source emmisions including incinerators, boilers and absorbers.

2.0 Equipment Required

- 2.1 GC/FID
- 2.2 GC/NDIR
- 2.3 Ice water bath: 1000ml Beaker filled with ice and deionized water.
- 2.4 Boiling water bath: 1000 ml Beaker filled with deionized water, wire gauze, tripod and Bunsen burner.
- 2.5 Dry ice.
- 2.6 Evacuated 1.8 liter vessel.
- 2.7 Condensate trap

3.0 Reagents Required

See 3.1 and 3.2 of SOP titled TCA 25.2/SOP

4.0 Procedure

This method requires the use of two different instruments, GC/FID and GC/NDIR and Orsat equipment. The samples are collected in two fractions, a gaseous fraction and a condensible fraction. Each fraction must be analyzed separately using both instruments.

- 4.1 Preparation of Gaseous Fraction of Sample
 - 4.1.1 Record room temperature and barometric pressure.
 - 4.1.2 When the sample has equilibrated to room temperature, measure the absolute pressure in the sample cylinder with a mercury manometer, and record the result as residual vacuum (R.V.).
 - 4.1.3 Pressurize the sample cylinder with ultra-pure nitrogen to at least 6 inches of mercury (absolute), then allow to equilibrate to room temperature (about 5 minutes).
 - 4.1.4 Repeat the measurement of absolute pressure, and record the result as pressure of nitrogen (PN₂).
 - 4.1.5 Calculation for dilution factor (XF) and sample amount (SA) (see 5.1).
- 4.2 Preparation of Condensable Fraction of Sample

Note: Inlet traps and outlet traps differ substantially in the level of hydrocarbons present, and so require slightly different handling. Several of the steps below will contain alternative instructions for onlet and outlet traps. Inlet traps and outlet traps are kept separate; inlet traps are not used to sample outlets, and vice versa in order to avoid sample carry-over problems.

- 4.2.1 Evacuate two 2 liter tanks to -30" Hg. Prepare a solution of 0.1N H₂SO₂ for the knock-out which is connected between the catalyst and the NDIR. The water knock-out also requires an ice bath.
- 4.2.2 Determine the baseline for the NDIR. On range 3, the baseline will usually be between 10 and 15 at room temperature.

- 4.2.3 Remove the trap from the freezer. Inlet traps: immerse the trap in a dry ice bath, connect one side of the trap to the carrier gas, and the other to an evacuated tank. Outlet traps: connect one side of the trap to the carrier gas, and the other side to a cold finger which is immersed in a dry ice bath (the trap itself is allowed to warm to room temperature). The other side of the cold finger is connected to an evacuted tank.
- 4.2.4 Start the carrier gas flowing through the trap for 5 minutes, until the pressure the pressure in the tank is about -10" Hg. This tank is then analyzed and added to the sample collected in the trap.
- 4.2.5 Connect the trap to the catalyst column. Attach a second evacuated cylinder to the outlet of the NDIR. If a inlet is being burned, use an eight liter cylinder. If an inlet is being burned, use a two liter flask. Perform a leak check by turning on the carrier gas flow briefly. Watch the bubbles in the water knock-out. When the bubbles stop with the system pressurized and the carrier gas is turned off, there should be a complete equlibrium in the drop-out. The presence of water moving backwards in the drop-out signifies a leak in the trap or catylist. If bubbles continue to flow, there is a leak in the NDIR or tank connection.
- 4.2.6 Start the carrier gas flow, remove the dry ice bath and begin heating the trap with the burners. For inlet traps, set the supplemental oxygen flow to match the carrier gas flow. For outlet traps, burn the cold finger along with the trap: supplemental oxygen is not needed in most cases. Observe the reading on the NDIR. It should quickly begin to rise.
- 4.2.7 Continue burning the trap until the NDIR reading falls to near the blank value. The trap is allowed to cool to room temperature, and then is briefly heated again to determine whether all the condensable hydrocarbons have been driven off. If the NDIR reading does not increase during this final check the trap is clean. It is important that hydrocarbons will condense in the tubing and

that it must be burned out with the trap. Also any place there is a connector, it must be heated to make sure that the inside is cleaned.

- 4.2.8 The tank from the burn and the purge are pressurized and analyzed (see above for gas sample portion).
- 4.3 GC/FID Analysis of Samples
 - 4.3.1 Check carrier gas (UP He), if low, change the gas tank.
 - 4.3.2 Change the output from "test" to "1", change the range from "Bal" to "10".
 - 4.3.3 Increase He to 50 psi.
 - 4.3.4 Turn on H2 to 30 psi.
 - 4.3.5 Ignite the FID.
 - 4.3.6 Turn on Air to 20 psi.
 - 4.3.7 Increase He to 80 psi.
 - 4.3.8 Turn FID Zero Suppression on, and switch detector Output from "1" to "2".
 - 4.3.9 Let the instrument warm up for 20 minutes.
 - 4.3.10 Start the Maxima program of the Dynamic Solutions system, and set up the automatic data acquisition according to the manual Maxima 820.
 - 4.3.11 Standard Calibration Calibration standards are commercial stock standard mixtures made from certified gas company.
 - (a) Inject a standard sample through a sample loop on the GC.
 - (b) Press "Run" on the GC to place the loop into the carrier gas circuit.
 - (c) The calibration standards are run in replicate until +/- 2% maximum deviation is obtained.
 - 4.3.12 The sample is injected in the same manner as the standard, steps 4.3.11: a, b, and c.

- 4.4 GC/NDIR Analysis of Samples
 - 4.4.1 Turn on the instrument and allow the GC oven to reach analysis temperature (approximately about 800 F).
 - 4.4.2 Start Nitrogen flow.
 - 4.4.3 Turn on the integrator: change the RANGE from "Tune" to "1", set GAIN from "0" to "5".
 - 4.4.4 When the GC oven reaches 800 F, turn the ZERO tune all the way to the right, and then adjust the SOURCE BALABCE until it reads between 35-40.
 - 4.4.5 Adjust the ZERO tune until the display reads "2" by turning it to the left.
 - 4.4.6 Equilibrate the Tenax loop in the ice water bath.
 - 4.4.7 Run a blank sample to check for carry-over prior to running standard.
 - 4.4.8 Standard Calibration Calibration standards are commercial stock standard from certified gas company.
 - (a) Flush the sample loop with the pressurized standard, then switch the injection valve to inject the sample.
 - (b) Press "Start" on the integrator.
 - (c) When the C_2 has been eluted or their expected retention time has passed, the carrier gas flow direction is reversed, and the Tenax loop is immersed in boiling water to desorb the remaining hydrocarbons.
 - (d) The calibration standards are repeated until +/- 5% maximum deviation is obtained.
 - 4.4.9 Inject the sample in the same manner as the standard, steps 4.4.8: a, b, and c
 - 4.4.10 Repeat the analysis of samples until +/-5% maximum deviation is obtained. If any of the hydrocarbons peaks are off scale, recalibrate the GC using a new standard that is closer to the concentration of the sample and then reanalyze the sample.

4.5 Dilution

See 4.5 of the SOP titled TCA 25.2

4.6 Orsat analysis of samples

See SOP titled 7.1

5.0 Calculations

- 5.1 Condensable Hydrocarbons
 - The intergrated area for the standard and the sample are available directly from the chart recorder.
 - Calculate the concentration in ppm of carbon 5.1.2 equivalent as follows:

$$C_{smp1} = \frac{C_{std} \times A_{smp1} \times V_{ves} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}{A_{std} \times V_{tank} \times \frac{P_{tank}}{29.9} \times \frac{520}{460 + T}}$$

where

 $C_{smpl} = concentration of the sample in ppm,$

C_{std} = concentration of the standard in ppm,

 A_{smpl} = area of the sample,

A_{std} = area of the standard,
V_{tank} = volume of the sample tank in liter,
V_{ves} = volume of the collection vessel in liter,
D_{ves} = neoggypo (checklists) = forth

P_{tank} = pressure (absolute) of the tank in inches of mercury,

= room temperature in F.

- Volatile hydrocarbons and gaseous components 5.2
 - The integrated area for the standard and the sample are available directly from the chart recorder.
 - 5.2.2 Calculate the concentration in ppm of carbon equivalent as follows:

$$C_{smpl} = \frac{C_{std} \times A_{smpl} \times \frac{(P_B + P_f)}{(P_B + P_i)}}{A_{std}}$$

where:

 P_B = barometric pressure (net) in inches of mercury,

P_i = residual pressure of sample tank,

P_f = final pressure of sample tank after N2 addition.

The following formula is used for the computation of the emission rate in lb/hr carbon:

$$\frac{\textit{C} \times 12 \textit{lb/mole} \times \textit{Q}_{\textit{sd}}}{3.79 \times 10^8 \textit{ft}^3 / \textit{lb.mole}}$$

where:

Q_{sd} = flow rate in standard cubic feet per hour (dry).



APPENDIX D
CALIBRATION DATA

PACIFIC ENVIRONMENTAL SERVICES, INC. —



(909) 887-2571 FAX. (909) 887-0549

CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS RECERTIFICATION

Customer

PACIFIC ENVIRONMENT SER ATTN: STEVE HERNANDEZ 13100 BROOKS DRIVE BALDWIN PARK CA91706

Assav Laboratory Scott Specialty Gases 2600 Cajon Boulevard San Bernardino, CA 92411

Purchase Order 0640-102 Scott Project # 25837.001

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number

ALM027046

Certification Date

12-23-93

Acid Rain Exp.

Cylinder Pressure

2000PSIG

Previous Certification Dates

05-20-93 General Exp. 12-23-95

ANALYZED CYLINDER

Components NITRIC OXIDE **Certified Concentration**

43.83PPM

Analytical Uncertainty* ±1% NIST Traceable

Balance Gas: Nitrogen

NOX

44.17PPM

*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

REFERENCE STANDARD

Type * GMIS :

Expiration Date

09-94

Cylinder Number

ALM033883

Concentration

100.4PPM

INSTRUMENTATION

Instrument/Model/Serial # TECO / 10AR-38644-258

Last Date Calibrated

10-29-93

Analytical Principle Chemi-Luminescent

ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

Components NITRIC OXIDE

ا والعمار بياو ميورين

Previous Certification

Date: 05-20-93	Respo	mse Units:mv
Z1=	R1=	T1=
R2=	72=	T2=
73=	T3=	R3=
Ave. Conc. of C	ust CvL	43.83PPM

Third Triad Analysis

Date:	12-23-93	R	espons	e Units: mv
Z1=	0.00	R1=	97.0	T1 = 41.8
R2=	97.0	Z2=	0.00	T2 = 41.8
Z3=	0.00	T3=	41.8	R3 = 97.0
Avg. Conc. of Cust Cyl.			43.83PPM	

Calibration Curve

Concentration= A =1.0083945	Ax+B	
B =-0.139473		

Date:	Response Units:			
Z1=	R1=	T1=		
R2=	72=	T2=		
Date: Z1= R2= Z3=	T3=	R3=		
Avg. Conc.	of Cust Cyl.			

Dute:	Response Units:		
Z1=	R1=	T1=	
R2=	Z2 =	T2=	
Dute: Z1= R2= Z3=	T3=	R3=	
Avg. Conc.	of Cust Cyl.		

Concentration=				
,				

Date:	Response Units:		
Z1=	R1=	T1=	
R2= Z3=	72=	T2=	
Z3=	T3=	R3=	
	of Coast Cal		

Date:	Response Units:		
Z1=	R1=	T1=	
Z1= R2=	Z2=	T2=	
23=	T3=	R3=	
Aug Conc	of Coat Cal		

Concentration=	

SPECIAL NOTES: IF THIS PRODUCT IS USED FOR ACID RAIN COMPLIANCE, THE ACID RAIN DATE NOTED ABOVE APPLIES PER 40 CFT PARY TO APPENDIX H. OTHERWISE THE GENERAL EXPIRATION DATE APLLIES.

Analyst



2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS

Pacific Environmental Services 13100 Brooks Drive Baldwin Park, CA 91706-0740

Assay Laboratory

Scott Specialty Gases 2600 Cajon Boulevard San Bernardino, CA 92411 Purchase Order 0640-102 Scott Project # 24700

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number

ALM027851

Certification Date

03-17-93

Expiration Date

Cylinder Pressure

1950 psig

Previous Certification Dates

NONE

09-13-94

ANALYZED CYLINDER

Components

NItric Oxide

Certified Concentration

22.51 ppm

Analytical Uncertainty* ± 1 % NIST Traceable

NOX

ict in

22.82 ppm

Balance Gas: Nitrogen

*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

REFERENCE STANDARD

Type: **SRM** 2629A **Expiration Date**

Cylinder Number FF28519

Concentration

19.3 ppm

INSTRUMENTATION Instrument/Model/Serial # TECO/ 10AR / 14853-150

Last Date Calibrated

1-20-93

Analytical Principle Chemi-Luminescent

Date:

Z1=

R2=

ANALYZER READINGS (Z=Zero Gas R=Reference Gas T=Test Gas r=Correlation Coefficient)

Date: Z1=

R2=

7.3=

Components

First Triad Analysis

Nitric Oxide

Date: 03-01-93 Response Units: mv				
Z1 = 0.000	R1=	78.33	T1=	90.95
R2= 78.30	Z2=	0.000	T2=	90.92
Z3= 0.000	T3-	90.94	R3=	78.28
Ave Cope of Cust Cvl. 22.41 pom				

Second Triad Analysis

Date: 03-17-93 Response Units: mv					nv
Z1=	0.000	R1=	77.12	T1=	90.20
R2=	77.24	Z2=	0.000	T2=	90.54
Z3 -	0.000	T3=	90.50	R3=	<i>7</i> 7.21
Avg. Conc. of Cust Cyl. 22.60 ppm					pm

R1=

Z2=

T3=

Response Units: mv

T1=

T2=

R3=

Calibration Curve

Concentration=	Ax ² +Bx+C	
A =-0.000504296		
B=1.0489880		
C =-0.292506		

Date:	Respo	onse Units: mv
Z1=	R1=	T1=
R2=	Z2 =	T2=
Z3=	T3=	R3=
Date: Z1= R2= Z3= Avg. Conc.	of Cust Cyl.	

R1= 72=

T3=

Avg. Conc. of Cust Cyl.

Respo	onse Units:	Date:	Date: Response Units	
=	T1=	Z1=	R1=	T1=
=	T2=	R2=	Z2 =	T2=
=	R3=	Z3=	T3=	R3=
Cyl.		Avg. Conc.	of Cust Cyl.	

Avg. Conc. of Cust Cyl.

Concentration=
,

Concentration=		

Special Notes



2600 CAJON BOULEVARD, SAN BERNARDINO, CA 92411

(909) 887-2571 FAX: (909) 887-0549

ACID RAIN DATE

CERTIFICATE OF ANALYSIS: EPA PROTOCOL GAS

PACIFIC ENVIRONMENTAL SER. PO# 0640-102 13100 BROOKS AVE

BALDWIN PARK, CA 91706

Assay Laboratory Scott Specialty Gases 2600 Caion Boulevard San Bernardino, CA 92411

Purchase Order 0640-102 27964.001 Project #

ANALYTICAL INFORMATION

Certified to exceed the minimum specifications of EPA Protocol 1 Procedure #G1, Section Number 3.0.4

Cylinder Number

ALM034155

Certification Date

10-06-93

GENERAL Date 10-06-95

Cylinder Pressure 1900 psig

ANALYZED CYLINDER

Components NITRIC OXIDE Certified Concentration

22.90 PPM

Analytical Uncertainty*

±1 % NIST Traceable

Balance Gas: Nitrogen

NOX

200 mark Silver

23.46 PPM

*Analytical uncertainty is inclusive of usual known error sources which at least includes reference standard error & precision of the measurement processes

REFERENCE STANDARD

Type --GMIS:

Expiration Date

Cylinder Number

ALM033911

Concentration

· 24.97 ppm

INSTRUMENTATION

Instrument/Model/Serial #

TECO / 10AR / 38644-258

Last Date Calibrated

07-26-93

Analytical Principle Chemi-Luminescent

ANALYZER READINGS (Z-Zero Gas R-Reference Gas T-Test Gas r-Correlation Coefficient)

Components

Nitric Oxide

First Triad Analysis

Second Triad Analysis

Calibration Curve

Z1 = 0.00R2= 96.4

Date: 09-29-93 Response Units: mv R1= 96.4 T1= 88.5 T2 = 88.572 = 0.00T3- 88.5 R3= 96.4 23 = 0.00Avg. Conc. of Cust Cyl. 22.91 ppm

Date: 10-06-93 Response Units: mv Z1= 0.00 R1= 96.4 T1= 88.4 T2= 88.3 R2= 96.4 72 = 0.00**Z3**-0.00 T3= 88.3 R3- 96.4 Avg. Conc. of Cust Cyl. 22.88 ppm

Concentration= Ax + B A =1.001 B=0.05283

Date:	Respo	onse Units: mv
Z1= R2=	R1=	T1=
R2=	22 =	T2=
23=	T3=	R3==
Avg. Conc	of Cust Cyl.	

Date:	Response Units: mv			
Z1=	R1=	T1=		
R2=	Z2=	T2=		
Z3=	T3=	R3=		
Avg. Conc. of Cust Cyl.				

Concentration=	
,	

Date:	Response Units: m		
Z1=	R1=	T1=	
R2=	Z2 =	T2=	
Z3=	T3=	R3=	
Avg. Conc.	of Cust Cyl.		

Dute: Z1= R2= Z3=	Response Units: mv		
Z1=	R1=	T1=	
R2=	Z2=	T2=	
Z3 =	T3=	R3=	
	of Cust Cyl.		

SPECIAL NOTES: IF THIS PRODUCT IS USED FOR ACID RAIN COMPLIANCE THE ACID RAIN EXPIRATION DATE NOTED ABOVE APPLIES PER 40 CFT PART 75,APPENDIX H. OTHERWISE THE GENERAL EXPIRATION DATE APPLIES. Thousas Wilson



Shipped From:

2600 CAJON BLVD.

SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

CERTIFICATE OF ANALYSIS

PACIFIC ENVIRONMENTAL SER

PO# 0640-102

13100 BROOKS DRIVE

PROJECT #: 02-28622-001

PO#: 0640-102

ITEM #: 02024520 4AL

DATE:11/11/93

BALDWIN PARK

CA 91706

CYLINDER #: ALMO35757

ANALYTICAL ACCURACY: +/-1%NIST

FILL PRESSURE: 2000PSIG

BLEND TYPE : CERTIFIED MASTER GAS

KERUES	LED GAS	HNHLIS	272
CONC_MOLES_		<u>(MOLES)</u>	
11.	PCT	11.09	PCT
50.	FFM	50.11	FIFM
11.	PCT	11.00	FCT
	BAL		BAL
		11. PCT 50. PPM 11. PCT	CONC MOLES (MOLES 11. PCT 11.09 50.11 11. PCT 11.00

2000PSIG BIN#2 11-12-93 CRM1675 14.08%CO2 ALMO01136 CRM1678 47.2PPM AAL5970 CO CRM2659 20.63% ALMO17555 D2

ANALYST:/

PLUMSTEADVILLE, PENNSYLVANIA / TROY. MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO BATON ROUGE, LOUISIANA



Shipped From:

2600 CAJON BLVD. SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

CERTIFICATE 0 F ANALYSIS

PACIFIC ENVIRONMENTAL SER

STEVE HERNANDEZ 13100 BROOKS DRIVE PROJECT #: 02-27898-001

FCT

FFM

PCT

BAL.

PO#: 0640-102

ITEM #: 02024520 2AL

DATE: 10/01/93

BALDWIN PARK

CA 91706

CYLINDER #: ALMO36879

ANALYTICAL ACCURACY: +-1%

FILL PRESSURE: 2000 PSIG

BLEND TYPE : ACUBLEND MASTER GAS

COMPONENT CARBON DIOXIDE CARBON MONOXIDE OXYGEN NITROGEN

REQUESTED GAS **ANALYSIS** CONC MOLES (MOLES) 18.00 18. PCT 75.00 75. F'F'M 19.00 19. P'CT BAL

2000 PSIG BIN#2 10-08-93 CRM1679 ALM10524 CO

CRM1675 ALMOO1136 CO2, CRM2659 ALMO17555 02

ANALYST:

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO BATON ROUGE, LOUISIANA



Shipped From: 2600 CAJON BLVD.

SAN BERNARDINO

CA 92411

Phone: 909-887-2571

Fax: 909-887-0549

CERTIFICATE OF ANALYSIS

PACIFIC ENVIRONMENTAL

STEVE HERNANDEZ

C/O FEDERAL EXPRESS

520 LAWERNCE EXPRESS WAY

SUNNYVALE

CA 94086

PROJECT #: 02-28936-001

PO#: F028-000

ITEM #: 02022913 4

DATE: 12/02/93

CYLINDER #: ALMO10841

ANALYTICAL ACCURACY: +/-1%NIST

FILL PRESSURE: 2000 PSIG

BLEND TYPE : CERTIFIED MASTER GAS

COMPONENT NITRIC OXIDE NITROGEN - OXYGEN FREE CONC MOLES

10. PPM
BAL

ANALYSIS
(MOLES)
10.36 PPM

BAL

NOX

10.39 PPM

2000 PSIG BIN #1 12-01 GRAVIMETRIC MASTER GAS AGAINST NIST CERTIFIED WEIGHT INDEPENDENT LABORATORY

CERTIFIED TO HAVE BEEN BLENDED AND VERIFIED TO BE CORRECT BY ANALYSIS.

ANALYST:

PLUMSTEADVILLE, PENNSYLVANIA / TROY, MICHIGAN / HOUSTON, TEXAS / DURHAM, NORTH CAROLINA SOUTH PLAINFIELD, NEW JERSEY / FREMONT, CALIFORNIA / WAKEFIELD, MASSACHUSETTS / LONGMONT, COLORADO BATON ROUGE, LOUISIANA EVER READY THERMOMETER CO., INC. 401 PARK AVENUE SOUTH NEW YORK, NY 10016

REPORT OF CALIBRATION

LIQUID-IN-GLASS-THERMOMETER

CALIBRATED BY EVER READY THERMOMETER CO.

MARKED: ERTCO 611-3FC S/N-2269

RANGE: -20 TO +110 DEGREES C IN 1 DEGREE GRADUATIONS.

THERMOMETER CORRECTION
READING (ITS-90)**

0.0 C 0.0 C 37.0 -0.1 56.0 0.0

** ALL TEMPERATURES IN THIS REPORT ARE BASED ON THE INTERNATIONAL TEMPERATURE SCALE OF 1990 (ITS-90) PUBLISHED IN THE METROLGIA 27, NO. 1, 3/10/90.

THIS THERMOMETER WAS CALIBRATED AGAINST A STANDARD CALIBRATED AT THE NATIONAL INSTITUTE OF STANDARDS AND TECHNOLOGY (NIST) FORMERLY THE NATIONAL BUREAU OF STANDARDS (NBS).

FOR A DISCUSION OF ACCURACIES ATTAINABLE WITH SUCH THERMOMETERS SEE NBS MONOGRAPH 150.

IF NO SIGN IS GIVEN ON THE CORRECTION, THE TRUE TEMPERATURE IS HIGHER THAN THE INDICATED TEMPERATURE; IF THE SIGN GIVEN IS NEGATIVE, THE TRUE TEMPERATURE IS LOWER THAN THE INDICATED TEMPERATURE. TO USE THE CORRECTIONS PROPERLY, REFERENCE SHOULD BE MADE TO THE NOTES GIVEN BELOW.

THE THERMOMETER WAS TESTED IN A LARGE, CLOSED-TOP, ELECTRICALLY HEATED, LIQUID BATH, BEING "IMMERSED" 76MM. THE TEMPERATURE OF THE ROOM WAS ABOUT 25 DEGREES C (77 DEGREES F). IF THE THERMOMETER IS USED UNDER CONDITIONS WHICH WOULD CAUSE THE AVERAGE TEMPERATURE OF THE EMERGENT LIQUID COLUMN TO DIFFER MARKEDLY FROM THAT PREVAILING IN THE TEST, APPRECIABLE DIFFERENCES IN THE INDICATIONS OF THE THERMOMETER WOULD RESULT.

THE TABULATED CORRECTIONS APPLY PROVIDED THE ICE POINT READING IS 0.0 DEGREES C. IF THE ICE-POINT READING IS FOUND TO BE HIGHER (OR LOWER) THAN STATED, ALL OTHER READINGS WILL BE HIGHER (OR LOWER) TO THE SAME EXTENT.

TEST NUMBER: 140381 DATE: 06/21/90

STANDARD SERIAL NO. 128239 NIST IDENTIFICATION NO. 88024 EVER READY THERMOMETER CO.



METER BOX TEMPERATURE READOUT CALIBRATION

DATE: 06-16-93

	Inlet (°F)	Outlet (°F)	Thermometer (°F)
BOX 1A	32	32	32 (0°C)
	77	76	77 (25°C)
	212	211	212 (100°C)
BOX 2A	32	32	32 (0°C)
	76	76	77 (25°C)
	213	213	212 (100°C)
BOX 3A	28	27	32 (0°C)
	75	75	77 (25°C)
	213	213	212 (100°C)

NOTE:

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. Number: 88024

PACIFIC ENVIRONMENTAL SERVICES, INC. —



THERMOCOUPLE CALIBRATION

CALIBRATED BY: Robert Aguner

DATE: 06-16-93

Thermocouple number	Thermocouple reading (°C)	Thermometer reading (°C)
TC-1	0.0 26.5 100.0	0.0 26.5 100.0
TC-2	OUT OF SERV	ICE
TC-3	0.0 27.5 99.0	0.0 26.5 100.0
TC-4	OUT OF SERV	ICE
TC-5	0.0 26.6 99.0	0.0 26.5 100.0
TC-6	0.0 26.5 100.0	0.0 26.0 100.0
TC-7	0.0 26.2 100.0	0.0 26.0 100.0
TC-8	0.0 26.3 100.0	0.0 26.0 100.0
TC-9	0.0 26.6 100.0	0.0 26.0 100.0

PACIFIC ENVIRONMENTAL SERVICES, INC. —

dia.	70.00	C 27 2
- 0 0		
\Box		

TC-10	0.0 26.3 100.0	0.0 26.0 100.0
TC-11	0.0 26.0 100.0	0.0 26.0 100.0
TC-12	0.0 26.0 100.5	0.0 25.0 100.0
C-1	NOT AVAILAE	3LE
C-2	NOT AVAILAE	BLE
S-1A	0.0 25.8 99.0	0.0 26.0 100.0
S-2A	0.0 25.5 99.0	0.0 26.0 100.0
S-16A	0.0 25.7 99.0	0.0 26.0 100.0
S-17A	0.0 25.9 99.0	0.0 26.0 100.0

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. Number: 88024

PACIFIC ENVIRONMENTAL SERVICES, INC.



DIGITAL THERMOMETER CALIBRATION

DATE: 06-16-93

Digital Thermometer	(°F)	Thermometer (°C)
Fisher	32.0 80.0 210.0	0.0 (32 °F) 26.5 (80 °F) 99.0 (210 °F)
Digital Thermometer	(°C)	Thermometer (°C)
Omega	1.0 26.0 98.0	0.0 26.5 98.0
Digital Thermometer	(°C)	Thermometer (°C)
Fluke (T1)	1.0 27.5 99.0	0.0 26.5 98.0
Digital Thermometer	(°C)	Thermometer (°C)
Fluke (T2)	1.1 27.4 99.0	0.0 28.0 98.0

NOTE:

Thermometer Standard Serial Number: 128239

Thermometer NIST I.D. 88024

PACIFIC ENVIRONMENTAL SERVICES, INC. —

DATE: DRY GAS METER IDENTIFICATION: 25507 3-31-93

BAROMETRIC PRESSURE (Pb):__

29,55

in. Hg

Model 5-190 OPERATOR WIN IDENTIFICATION: AL-20

PACIFIC ENVIRONMENTAL SERVICES, INC.

$Q = 17.65 \frac{V_1 - P_1}{\Theta (I_1 + 460)}$	C	D C	1.20 🗚	C	Oc.	1.00 4	٠ .	Pb: 29.46 + B	0.80 A	C	P6 = 29.48 - B	0.60 1	C	<i>t</i> 3	0.40 A		cfm	APPROXIMATE	(Note: rb
۳,	11.100	_	L					10,130	10,122	7.435	7.425	7.440	S. 392	5.392	5.420	ft³	GAS VOLUME	MET TEST	e was seemed
$= \frac{V_{\star}}{V_{dg}} \times \frac{(\bar{I}_d + 460)}{(\bar{I}_s + 460)} \times$	414.306	852.414	402,758	390,237	369, 142	368, 410	356.529	335,979	325.721	316.689	316,639	301.605	300,968	290,028	284.603	;	(V _{dg})	DRY GAS	Comme Person
$(P_b + \frac{AP}{13.6})$	74	24	75	74 74	74 74	74 74	74 74	74 74	74 74	77	74 74	74 74	74 74	74 74	74 74	αρ •F	WET TEST		
	1	1	,	1	1	,	,	,	1	ſ	1	1	1	l	1	INLET (T _i) °F		TEMPER	1
	18.4	78, 2 78, 2	78,2	78.2	78,2	78,2	78, 2 78, 2	78,2	78, Z 78, Z	77.4	77.4	77.4	77.4	77.4	77.1	OUTLET (T _o) °F	DRY GAS METER	TEMPERATURES	
	78	78	38	78	78	78	78	78	78	78	77	77	77	77	77	AVERAGE (T _d) °F	ER		
	-7.10	-7.10	-7,10	-5.15	-21.72	-212 -212	-3.65	-3,65	-3.65	-2.35	- 2.35	-2,35	-1,35	-1.35	7.35	in H ₂ 0	PRESSURE (AP)	DRY GAS	
	9.0	9.0	9,0	10.0	10.0	10.0	12.0	12.0	12.0	12.0	12.0	12.0	13.0	13.0	13.0		min.	TIME	
	1,201	1.204	1,203	1.005	1.007	1.008	0,825	0.812	0.822	0.604	0,603	0.606	0,405	0,405	0.407		cfm	FLOW RATE	
	1.007	1.007	1,008	1.009	1.009	1.009	1.009	1.010	1.010	1.012	1.009	1.010	1.009	800%	1.008		COEFFICIENT	METER	
	1,007	1	1	1.009		1	1.010		•	1.010			1.008				(Y ₀ g)	AVERAGE METER	

DRY GAS METER AND ORIFICE CALIBRATION

Barometric Pressure		Dry Gas Meter No.
29.8	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	1557640
Calibration Date		Meter Box No. 3A
12-28-93	5 6 9 9 1 1 1	¥
Calibrated by:		Reference Dry G
		Reference Dry Gas Meter No. 25507 Rockwell
	113	e 29.8 Calibration Date 12-28-93 Calibrated by:

4 9.4 1 1.05 1 0.991 1 1.931	9.4 [1.05 [0.99 [1.93]	**************************************	74	87	101	74	10.376	741.358 I	730.982 I	10.01	802.990 I	1 792,980 I	4.0
1.0	13.4 0.74 1.003 1.968	13.4 [92	85 1	97	74	10.260	730.205 I	719.945 [10.023 [792.290 I	782.267	2.0
0.99	7 18.7 0.53 0.994 1.903	18.7 [87 [1 82 1	I 92	7.6	10.378	719.423 [709.045 I	10.132 [781.789 I	771.657	1.0
0.99	25.6 0.38 0.998 1.837	25,6 [84 [92 1	9.6	76	10.167	708.743 [698.576	10,011	771.359	761.348	0.5
He He	G=cfm		Hox Dun Average	He He He He He He He He		Reference	Volume Vd=ft3	Reading I V=ft3 I	Reading I V=ft3 I	Volume I Vr=ft3 I	Reading I V=ft3 I	DGM Reading V=ft3	Manometer Setting H="H2O
		· Ru				[]	Test	Final I	Initial I	Reference I	G.	Ē	Orifice
				ture	Temperature			provide		a	-	-	

	AVERAGE		
	-	p-m-+ 2	
	0.997	ber wel	
-	1,910		
-	-	perced	brene



MAGNEHELIC CALIBRATION CHECK

	LOW		MEI		HIC	
<u>UNIT</u>	Mag	Man	Mag	Man	Mag	Man
05-24-93						
JW16 (0-1")	0.10	0.10	0.47	0.47	0.90	0.90
GF17 (0-0.5")	0.060	0.060	0.260	0.260	0.470	0.485
CA59 (0-4")	0.50	0.50	1.70	1.70	3.50	3.50
07-30-93						
R15E (0-1")	0.12	0.12	0.49	0.49	0.82	0.82
JW16 (0-1")	0.12	0.12	0.50	0.50	0.82	0.82
GF17 (0-0.5")	0.065	0.060	0.260	0.260	0.430	0.420
CA59 (0-4")	0.40	0.38	1.90	1.90	3.40	3.40
11-26-93						
R15E (0-1")	0.11	0.10	0.51	0.51	0.87	0.87
JW16 (0-1")	0.10	0.10	0.51	0.51	0.89	0.89
GF17 (0-0.5")	0.050	0.045	0.250	0.250	0.430	0.440
CA59 (0-4")	0.51	0.49	2.02	2.00	3.65	3.65

Readings in " $\rm H_2O$

Reference: Oil Manometer - Dwyer #400-10 0-1" inclined, 1-10" vertical

PACIFIC ENVIRONMENTAL SERVICES, INC.



PRECISION INSTRUMENT REPAIR CO.

13414 WOODRUFF AVE., BELLFLOWER, CA. 90706

310 /925-6672

REPORT NO: 3916

Certification Report

Precision Balances and Scales

This is to certify that the balance calibrated is in compliance to US GOVERNMENT MILITARY BOOKLET, MIL-STD-45662A and that the standards used meet the compliance of NIST (NATIONAL INSTITUTE OF STANDARDS AND TECHNOLOGY)

PRODUCT ORIGINAL CONDITION: In working order.
X CALIBRATED AT IN-USE LOCATION CALIBRATED AT P.I.R. OFFICE.
TEMPERATURE: 70°F HUMIDITY: 59% STANDARD USED: Class I S/N 5943

MANUFACTURER, MODEL, SERIAL NUMBER OF PRODUCT CALIBRATED

1 TORSION TORBAL EA-1 ANALYTICAL BALANCE, S/N 156636

Calibrated to: 100mg=100.0mg

Linearity: OK Maximum Load: OK Repeteability: OK

COMPANY OR PRODUCT OWNER:

PACIFIC ENVIRONMENTAL SERVICES 13100 BROOKS, SUITE 100 BALDWIN PARK, CA. 91706 ATTN: SIYA

PO# 0640-102



FI

Date of Calibration 12-21-93 Technichian Tom Benson License 1-0903



HYDROCARBON CYLINDERS VOLUME CALIBRATION 08-20-90

Cylinder #	Full, lbs	Tare, lbs	Net, 1bs	Volume, L
101	31.33	4.45	26.88	12.22
102	32.65	5.83	26.82	12.19
103	31.60	4.69	26.91	12.23
104	31.65	4.68	26.97	12.26
105	31.42	4.47	26.95	12.25
106	32.62	5.82	26.80	12.18
107	31.14	4.50	26.64	12.11
108	32.45	5.98	26.47	12.03
109	31.22	4.69	26.53	12.06
110	31.15	4.68	26.47	12.03
Supplemental	Set 07-29-92			
111	31.42	4.86	26.56	12.07
112	32.44	5.86	26.58	12.08
113	32.44	5.96	26.48	12.03
114	31.50	4.48	27.02	12.28
115	32.45	5.90	26.55	12.06
116	31.55	4.45	27.10	12.31
117	31.35	4.71	26.64	12.10
118	32.62	6.04	26.58	12.08
119	31.40	4.73	26.67	12.12
120	31.50	4.78	26.72	12.14

Note: Cylinders were filled to the rim (not including fittings) with water and weighed on a 125-lb capacity platform balance.

Water density at room temperature = 0.997 gm/cc or 0.0022 lbs/cc.



Pitot Tube Calibration Data Sheet

Calibrated by: Anga Mish	<u> </u>		_
Date: 12-30-93			_
Pitot Tube I.D. S-17 A			_
Effective Length: 5'			_
Pitot Tube Assembly Level ?	Yes	□ No	
Pitot Tube Openings Damaged ?	☐ Yes	□ No	
If Yes, Explain			_
			_
Δ- a			
$\alpha_1 = \underline{\qquad \qquad }^{\circ}(<10^{\circ})$	$\alpha_2 =$	°(<10°)	
$\beta_1 = $	$\beta_2 =$	°(<10°)	
γ = ο θ =	0 •	A = 0.954	,
$Z = A \sin \gamma = $	cm (in.)	0.32 cm (<% in.)	
$W = A \sin \theta = \underline{\hspace{1cm}}$	cm (in.)	0.08 cm (<1/32 in.)	
$P_A = $	cm (in.)		
$P_{B} = 0.477$	_ cm (in)		
$D_t = \underline{0.375}$	_ cm (in.)		
Comments:			
		· · · · · · · · · · · · · · · · · · ·	
Calibration Required ?	☐ Yes	□ No	

- PACIFIC ENVIRONMENTAL SERVICES. INC. -